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18: gb_pa2.*
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20:  em_hgtgo_hum.*
21:  em_hgtgo_inv.*
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42:  em_in.*
43:  em_or.*

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SUMMARIES

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3	1637	51.5	2144	88	AF051426	Homo sapi
4	1227	38.6	104123	93	HSAC6345.3	Continuation (4 of
5	1176	37.0	137932	86	AC005950	Homo sapi
6	1176	37.0	181483	63	AC013791	Homo sapi
7	1074	33.8	244254	93	HSAC001128	Con
8	548	17.2	155074	85	AC003693	Human Chr

[illegible]

QY 715 gtgggaccttggggcgctgctgcttggccggaggccatttccatcatcgacctc 774
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QY 1975 aggttggcactcatcacgacatgcttccacagctgctctcttgcacaggtgagcagacc 2034
Db 1923 AGGTGGCAGCTCATCCGACATGCTTACCAGCTCTCTCTGACGGTGGCAGCACC 1982
QY 2035 cccggcagcgcgccccccagagagggcggtggccacatcacccagccctcgcgagct 2094
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DEFINITION Homo sapiens clone RP11-19N21, WORKING DRAFT SEQUENCE, 9 unordered
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ACCESSION AC013791
VERSION AC013791.4 GI:12313824
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 181483)
AUTHORS Birren,B., Linton,L., Nusbaum,C. and Lander,E.
TITLE Homo sapiens, clone RP11-19N21
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 181483)
AUTHORS Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,
Baldwin,J., Barna,N., Beckerly,R., Boguslavsky,L., Boukhgalter,B.,
Brown,A., Castle,A., Collangelo,M., Collins,S., Collymore,A.,
Cooke,P., DeArellano,K., Dewar,K., Domino,M., Donelan,L., Doyle,M.,
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Galagan,J., Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L.,
Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J.,
Lehocsky,J., Lieu,C., Locke,K., Macdonald,P., Marquis,N.,
McEwan,P., McGurk,A., McKernan,K., McLaughlin,J., Meldrum,J.,
Morrow,J., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,
Peterson,K., Pollara,V., Riley,R., Roy,A., Santos,R., Severy,P.,
Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,
Tesfaye,S., Tirrell,A., Vassiliev,H., Vo,A., Wheeler,J., Wu,X.,
Wyman,D., Ye,W.J., Zimmer,A. and Zody,M.
Direct Submission
Submitted (15-NOV-1999) Whitehead Institute/MIT Center for Genome
Research, 220 Charles Street, Cambridge, MA 02141, USA
On Jan 19, 2001 this sequence version replaced gi:7382100.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L3857
Center Clone name: 19_N.21
----- Summary Statistics
Sequencing vector: M13; M77815; 47% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 176802 bases at least Q40
Consensus quality: 179250 bases at least Q30
Consensus quality: 180075 bases at least Q20
Insert size: 173000; agarose-fp
Insert size: 180683; sum-of-contigs
Quality coverage: 10.2 in Q20 bases; agarose-fp
Quality coverage: 9.8 in Q20 b.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 9 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 35935: contig of 35935 bp in length
* 35936 36035: gap of 100 bp
* 36036 37309: contig of 1274 bp in length
* 37310 37409: gap of 100 bp
* 37410 38442: contig of 1033 bp in length
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* 63015 63114: gap of 100 bp
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[illegible]

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Matches 1224; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

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LOCUS Human Chromosome 11p15.5 PAC clone pDJ915f1 containing KvLQ1 gene,
DEFINITION complete sequence.
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VERSION AC003693.1 GI:3687269
KEYWORDS HTG.
SOURCE human.
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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 155074)
AUTHORS Evans,G.A., Athanasiou,M., Aguayo,P., Armstrong,D., Basit,M.,
Buetner,J., Bumeister,R., Card,P., desaliboat,F., Dunn,J.,
English,C., Ethridge,S., Garner,H.R., Gee,V., Gordon,M., Gotway,G.,
Grant,O., Hahner,L., Joslin,J., Lewis,E., Loo,H., Loo,K.N.,
Major,T., McFarland,J., Newton,J., Osborne-Lawrence,S.,
Schageman,J., Schultz,R.A., Stimson,S., Syed,M. and Ward,T.
HTG Submission
Unpublished
2 (bases 1 to 155074)
Evans,G.A., Athanasiou,M., Basit,M., Bradbury,P., Brignac,S.,
Bumesiter,R., Davis,C., English,C., Franklin,T.L., Garner,H.R.,
Gee,V., Gordon,M., Gotway,G., Grant,O., Hahner,L., Harris,J.,
Hinson,S., Narayanaswamy,U., Newton,J., O'Brien,K., Patel,P.,
Schageman,J., Schilling,P., Schultz,R., Syed,M., Valenzuela,D.,
Ward,T. and Wilson,R.
Direct Submission
Submitted (17-DEC-1997) Genome Science & Technology Center,
University of Texas Southwestern Medical Center, 5323 Harry Hines
Blvd, Dallas, TX 75235-8591, USA
3 (bases 1 to 155074)
Evans,G.A., Athanasiou,M., Aguayo,P., Armstrong,D., Basit,M.,
Buetner,J., Butler,C., Card,P., desaliboat,F., Dunn,J.,
English,C., Ethridge,S., Garner,H.R., Gee,V., Gordon,M., Gotway,G.,
Grant,O., Hahner,L., Joslin,J., Lewis,E., Loo,H., Loo,K.N.,
Major,T., McFarland,J., Newton,J., Osborne-Lawrence,S.,
Schageman,J., Schultz,R.A., Stimson,S., Waller,K. and Ward,T.
Direct Submission
Submitted (30-SEP-1998) Genome Science & Technology Center,
University of Texas Southwestern Medical Center, 5323 Harry Hines
Blvd, Dallas, TX 75235-8591, USA
On Oct 1, 1998 this sequence version replaced gi:3264564.
Further information regarding the map of this region or
annotation of pDJ915f1 can be found at
http://gestec.swmed.edu/chromosome.htm
IMPORTANT: This submission contains the entire insert of clone
pDJ915f1. pDJ915f1 comes from the RPCI-3 PAC library constructed
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KEYWORDS KVLQTL1.
SEGMENT 2 of 17
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ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
1 (sites)
Tanaka, T., Tanaka, T., Nagai, R., Kikuchi, K., Ogawa, S., Okada, S.,
Yamagata, S., Yano, K., Yazaki, Y. and Nakamura, Y.
Genomic organization and mutational analysis of KVLQTL1, a gene
responsible for familial long QT syndrome
Hum. Genet. 103 (3), 290-294 (1998)
JOURNAL 99013427
MEDLINE 2 (bases 1 to 471)
REFERENCE Tanaka, T.
AUTHORS Direct Submission
JOURNAL Submitted (01-JUN-1998) to the DDBJ/EMBL/GenBank databases.
Toshhiro Tanaka, Institute of Medical Science, University of
Tokyo, Laboratory of Molecular Medicine; 4-6-1 Shirokanedai,
Minato-ku, Tokyo 108-8639, Japan
(E-mail: toshitan@ims.u-tokyo.ac.jp, Tel: 81-3-5449-5374,
Fax: 81-3-5449-5406)
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ACCESSION U86146
VERSION U86146.1 GI:2076879
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
1 (bases 1 to 386)
Tanaka, W.P., Levesque, P.C., Little, W.A., Conder, M.L., Shalaby, F.Y.
and Blannar, M.A.
KVLQTL1, a voltage-gated potassium channel responsible for human
cardiac arrhythmias
Proc. Natl. Acad. Sci. U.S.A. 94 (8), 4017-4021 (1997)
JOURNAL 97268689
MEDLINE 2 (bases 1 to 386)
REFERENCE Yang, W.P., Levesque, P.C., Little, W.A., Conder, M.L., Shalaby, F.Y.
AUTHORS and Blannar, M.A.
Direct Submission
JOURNAL Submitted (21-JAN-1997) Cardiovascular Drug Discovery,
Bristol-Myers Squibb PRI, K14-01, P.O. Box 4000, Princeton, NJ
08543-4000, USA
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VERSION	ABO15153.1 GI:3953626
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SEGMENT	7 of 17
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ORGANISM	Homo sapiens
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REFERENCE AUTHORS	TITLE	JOURNAL MEDLINE REFERENCE AUTHORS	TITLE JOURNAL
1. HARRIS, R. J. & HARRIS, R. J.	THE EFFECT OF THE HARRIS, R. J. & HARRIS, R. J.	THE EFFECT OF THE HARRIS, R. J. & HARRIS, R. J.	THE EFFECT OF THE HARRIS, R. J. & HARRIS, R. J.

Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
1 (sites)
Itoh, T., Tanaka, T., Nagai, R., Kikuchi, K., Ogawa, S., Okada, S.,
Yamagata, S., Yano, K., Yazaki, Y., and Nakamura, Y.
Genomic organization and mutational analysis of KVLQT1, a gene
responsible for familial long QT syndrome
Hum. Genet. 103 (3), 290-294 (1998)
99013427
2 (bases 1 to 237)
Tanaka, T.
Direct Submission
Submitted (01-JUN-1998) to the DDBJ/EMBL/GenBank databases.
Toshinhiro Tanaka, Institute of Medical Science, University of
Tokyo, Laboratory of Molecular Medicine; 4-6-1 Shirokanedai,
Minato-ku, Tokyo 108-8639, Japan
(E-mail: toshitan@ims.u-tokyo.ac.jp, Tel: 81-3-5449-5374,
Fax: 81-3-5449-5406)

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; APPLICANT: Malsen, Gareth
; APPLICANT: Townley, David
; APPLICANT: Morris, MacDonald
; TITLE OF INVENTION: Single Nucleotide Polymorphisms Associated With ADME Genes
; FILE REFERENCE: GX-0013-5 P
; CURRENT APPLICATION NUMBER: US/60/313,371
; CURRENT FILING DATE: 2001-08-16
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; SOFTWARE: PERL Program
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; APPLICANT: Morris, MacDonald
; APPLICANT: Lal, Preeti
; APPLICANT: Diep, Dinh
; TITLE OF INVENTION: METHOD FOR THE IDENTIFICATION OF SEQUENCE POLYMORPHISMS USING
; TITLE OF INVENTION: POLYNUCLEOTIDE SEQUENCE DATABASES, AND SINGLE NUCLEOTIDE
; FILE REFERENCE: GX-0019-1 P
; CURRENT APPLICATION NUMBER: US/60/324,185
; CURRENT FILING DATE: 2001-09-21
; NUMBER OF SEQ ID NOS: 35862
; SOFTWARE: PERL Program
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; LENGTH: 4833
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; ORGANISM: Homo sapiens
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; NAME/KEY: misc_feature
; OTHER INFORMATION: Incyte ID No: 348127.7
US-60-324-185-25008
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Query Match 83.3%; Score 2651; DB 10; Length 4833;

Best Local Similarity 100.0%; Pred. No. 0;

Matches 2701; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Qy 480 caaggccgcgtctacaaacttctcagagctccacacggctggaaatgcttcttacc 539
Db 2116 caaggccgcgtctacaaacttctcagagctccacacggctggaaatgcttcttacc 2175
Qy 540 ctccgcgtcttctcagctgctgctccctcatcttcagcgtgcttccaccatcga 599
Db 2176 ctccgcgtcttctcagctgctgctccctcatcttcagcgtgcttccaccatcga 2235
Qy 600 gcagtagccgcctggccacgggagctctcttctgagtgagatcgtggtggtgtt 659
Db 2236 gcagtagccgcctggccacgggagctctcttctgagtgagatcgtggtggtgtt 2295
Qy 660 ctccggagcaggtacgtggtccgctctgttcccggaagccatttccatcatcgaggt 719
Db 2296 ctccggagcaggtacgtggtccgctctgttcccggaagccatttccatcatcgaggt 2355
Qy 720 cctctggggcgagctgcgttcttcccggaagccatttccatcatcgaggt 779
Db 2356 cctctggggcgagctgcgttcttcccggaagccatttccatcatcgaggt 2415
Qy 780 cgtggcctccatggtgctctcgtgggctcagagggcaggtgtttgccagctcgc 839
Db 2416 cgtggcctccatggtgctctcgtgggctcagagggcaggtgtttgccagctcgc 2475
Qy 840 catcaggggcactcgccttctcagatcctgaggtgctacagctgacgccaggaggt 899
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Qy 3060 ggcattacatgcagataagaaatcaataatttggtggtatttgatctgtgttttaagt 3119
|||
Db 4696 ggcattacatgcagataagaaatcaataatttggtggtatttgatctgtgttttaagt 4755

Qy 3120 ttacagtgattttgatttatttattgcaagcttttccataataaaacgtggagaatca 3179
|||
Db 4756 ttacagtgattttgatttatttattgcaagcttttccataataaaacgtggagaatca 4815

Qy 3180 ca 3181
||
Db 4816 ca 4817

RESULT 4
US-60-313-371-1500
; Sequence 1500, Application US/60313371
; GENERAL INFORMATION:
; APPLICANT: Ring, Huijun z.
; APPLICANT: Malsen, Gareth
; APPLICANT: Townley, David
; APPLICANT: Morris, MacDonald
; TITLE OF INVENTION: Single Nucleotide Polymorphisms Associated with ADME Genes
; FILE REFERENCE: GX-0013-5 P
; CURRENT APPLICATION NUMBER: US/60/313,371
; CURRENT FILING DATE: 2001-08-16
; NUMBER OF SEQ ID NOS: 2447
; SOFTWARE: PERL Program
; SEQ ID NO 1500
; LENGTH: 2702
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; OTHER INFORMATION: KCMQ1_mrna_build.1
US-60-313-371-1500
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Query Match 82.2%; Score 2615; DB 10; Length 2702;
Best Local Similarity 100.0%; Pred. No. 0;
Matches 2665; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
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Qy 484 ggcgcgtctcaactctctcagagctccaccgctggaaatgtctgttaccacttc 543
Db 37 ggcgcgtctcaactctctcagagctccaccgctggaaatgtctgttaccacttc 96

Qy 544 gccgtctctcctcatcgtcgtgctgcctcatcttcagcgtcgtctccaccatcgagcag 603
|||
Db 97 gccgtctctcctcatcgtcgtgctgcctcatcttcagcgtcgtctccaccatcgagcag 156

Qy 604 tatgcgccttgccacgggactctctcttgatggagatcgtggtgtttcttc 663
|||
Db 157 tatgcgccttgccacgggactctctcttgatggagatcgtggtgtttcttc 216

Qy 664 gggacggagtacgtggtcgcctctggtccgcgctgctccgcagcagtaagtgggcctc 723
|||
Db 217 gggacggagtacgtggtcgcctctggtccgcgctgctccgcagcagtaagtgggcctc 276

Qy 724 tggggggcgtgcgttttggccggaagccatttccatcgcacccatcgtggtcg 783
|||
Db 277 tggggggcgtgcgttttggccggaagccatttccatcgcacccatcgtggtcg 336

Qy 784 gccctccatgggtggtcctcgtggtggtcccaagggtgtttggccacgtcgccatc 843
|||
Db 337 gccctccatgggtggtcctcgtggtggtcccaagggtgtttggccacgtcgccatc 396

Qy 844 agggcaccgtctcctcagatcctgagatgctacacgtgcacccgcagggagggcacc 903
|||
Db 397 agggcaccgtctcctcagatcctgagatgctacacgtgcacccgcagggagggcacc 456

Qy 904 tggaggtccctgggctcgtggtgtcttcacaccgccagggaggtgataaccacctgtac 963
|||
Db 457 tggaggtccctgggctcgtggtgtcttcacaccgccagggaggtgataaccacctgtac 516
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Qy 964 atcggcttctcctggcctcatctctctcgtactttgttaccttggtgagaaagcgcg 1023
|||
Db 517 atcggcttctcctggcctcatctctctcgtactttgttaccttggtgagaaagcgcg 576

Qy 1024 gtgaacgagtcagggccgcgtggagttcggcagctacgcagatgcgtgtggtgggggtg 1083
|||
Db 577 gtgaacgagtcagggccgcgtggagttcggcagctacgcagatgcgtgtggtgggggtg 636

Qy 1084 gtcaacagtcaccaccatcggtctatggggacaaagtgccccacagcgtgggtcggaagacc 1143
|||
Db 637 gtcaacagtcaccaccatcggtctatggggacaaagtgccccacagcgtgggtcggaagacc 696

Qy 1144 atcgcctcctgcttctcgtcttggcatcctctcttcttgcgtccacggggattctt 1203
|||
Db 697 atcgcctcctgcttctcgtcttggcatcctctcttcttgcgtccacggggattctt 756

Qy 1204 ggcctcggggtttgccctgaaggtgcagcagaagcagagggcagaaacttcaaccggcag 1263
|||
Db 757 ggcctcggggtttgccctgaaggtgcagcagaagcagagggcagaaacttcaaccggcag 816

Qy 1264 atccccggcggcagcctcactcatctcagaccgcatggaggtgctatgctgcgagaacccc 1323
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Db 817 atccccggcggcagcctcactcatctcagaccgcatggaggtgctatgctgcgagaacccc 876

Qy 1324 gactcctcaactggaaagatctacatccggaaagccccccggagggcacaactctctgtca 1383
|||
Db 877 gactcctcaactggaaagatctacatccggaaagccccccggagggcacaactctctgtca 936

Qy 1384 ccagcccccaaccccaagaaagtctgtggttaaaagaaaaaagtccaagctggacaaa 1443
|||
Db 937 ccagcccccaaccccaagaaagtctgtggttaaaagaaaaaagtccaagctggacaaa 996

Qy 1444 gacaatggggtgactcctcctggagagaagatctcacagtcctcccatatcacgtgcgacccc 1503
|||
Db 997 gacaatggggtgactcctcctggagagaagatctcacagtcctcccatatcacgtgcgacccc 1056

Qy 1504 ccagaaagagcggcgtggaacactctctctgcagcgtctatgacagttctgtaagaag 1563
|||
Db 1057 ccagaaagagcggcgtggaacactctctctgcagcgtctatgacagttctgtaagaag 1116

Qy 1564 agcccaactcgtggaagtgcagtcgcccatttcatgagaccacacagcttcgcgag 1623
|||
Db 1117 agcccaactcgtggaagtgcagtcgcccatttcatgagaccacacagcttcgcgag 1176

Qy 1624 gacctggacctggaaggaggagactctgctgaccccatcacccatctcacagctcg 1683
|||
Db 1177 gacctggacctggaaggaggagactctgctgaccccatcacccatctcacagctcg 1236

Qy 1684 gaacacatcggggccaccattaaagggtatcttcagcagatcagtaactttgtggcagaag 1743
|||
Db 1237 gaacacatcggggccaccattaaagggtatcttcagcagatcagtaactttgtggcagaag 1296

Qy 1744 aaattccagaacgcggaagccttacgatgtcgggagcgtattgagcagactcgcag 1803
|||
Db 1297 aaattccagaacgcggaagccttacgatgtcgggagcgtattgagcagactcgcag 1356

Qy 1804 ggcacctcaacctcatggtgcgcatacaggagctgcagaggaggtggaccagtcatt 1863
|||
Db 1357 ggcacctcaacctcatggtgcgcatacaggagctgcagaggaggtggaccagtcatt 1416

Qy 1864 ggaagccctcactgttcatctcgtctcagaaaagagcagatcgcgagcaaacg 1923
|||
Db 1417 ggaagccctcactgttcatctcgtctcagaaaagagcagatcgcgagcaaacg 1476

Qy 1924 atcggcgcgcctcgaacagtagaagacaaggtgaacagctggaacagaggtgga 1983
|||
Db 1477 atcggcgcgcctcgaacagtagaagacaaggtgaacagctggaacagaggtgga 1536

Qy 1984 ctcatcacgacatgcttaccagctgctctcttgcagcgtggcagcccccgagc 2043
|||
Db 1537 ctcatcacgacatgcttaccagctgctctcttgcagcgtggcagcccccgagc 1596

Qy 2044 ggcggccccccccagagaggggcgggccccacataccccagccctgcggcgtcc 2103
|||
```


Qy	1144	atcgctccctgcttctctgtcttcttgccatctcttcttgctcccgagggaattctt	1201
Db	697	atcgctccctgcttctctgtcttggcatctcttcttgctcccgagggaattctt	756
Qy	1204	ggctcggggttgccctaagctgcgcagaagcaagcagaagcacttcaaccggcag	1263
Db	757	ggctcggggttgccctaagctgcgcagaagcagaagcacttcaaccggcag	816
Qy	1264	atcccgcgcgagctcactcattcagaccgatggaggtgctatgtgcccgaagaacccc	1323
Db	817	atcccgcgcgagctcactcattcagaccgatggaggtgctatgtgcccgaagaacccc	876
Qy	1324	gactctccactggaagatctacatccggaagcccccgagccacactctgctgtca	1383
Db	877	gactctccactggaagatctacatccggaagcccccgagccacactctgctgtca	936
Qy	1384	cccagccccaaaaccagaagctctgtgtggttaagaaaaaaagtctcaagctggacaaa	1443
Db	937	cccagccccaaaaccagaagctctgtgtggttaagaaaaaaagtctcaagctggacaaa	996
Qy	1444	gacaatgggggtgactccctggagagaagatgcttcacagtcctcccatatcacgtgcgacccc	1503
Db	997	gacaatgggggtgactccctggagagaagatgcttcacagtcctcccatatcacgtgcgacccc	1056
Qy	1504	ccagaagagcgcggtgagcaactctctgtcgacggctatgacagttctgttaaggaaag	1563
Db	1057	ccagaagagcgcggtgagcaactctctgtcgacggctatgacagttctgttaaggaaag	1116
Qy	1564	agcccaacactgctggaagtgagcatgccccatttcatgagaacbaacagttctgcgcgag	1623
Db	1117	agcccaacactgctggaagtgagcatgccccatttcatgagaacbaacagttctgcgcgag	1176
Qy	1624	gacctggacctggaaagggaagactctgtgacacccatcacccacatctcaagctgfcgg	1683
Db	1177	gacctggacctggaaagggaagactctgtgacacccatcacccacatctcaagctgfcgg	1236
Qy	1684	gaacacatcgggccacattaaagtcatctcgacgcatgcagttcttggcccagaag	1743
Db	1237	gaacacatcgggccacattaaagtcatctcgacgcatgcagttcttggcccagaag	1296
Qy	1744	aaattcagaagcgcggaagccttacgatgtgcgggagctcatttgagcagttactgcgag	1803
Db	1297	aaattcagaagcgcggaagccttacgatgtgcgggagctcatttgagcagttactgcgag	1356
Qy	1804	ggccacctcaacctcatggtgcgcatacaagagctgcagagagctgacacagttccatt	1863
Db	1357	ggccacctcaacctcatggtgcgcatacaagagctgcagagagctgacacagttccatt	1416
Qy	1864	gggaagccctcactgttcatctccgtctcagaataaagacaagatcgcgcgagcaacag	1923
Db	1417	gggaagccctcactgttcatctccgtctcagaataaagacaagatcgcgcgagcaacag	1476
Qy	1924	atcgggcccgccctgaaccgagtagaacaaggtgacgagctggaccagaggtcgga	1983
Db	1477	atcgggcccgccctgaaccgagtagaacaaggtgacgagctggaccagaggtcgga	1536
Qy	1984	ctcatccgcacatgcttcaaccagctgctctctcttgacgggtggcagaccccccgagc	2043
Db	1537	ctcatccgcacatgcttcaaccagctgctctctcttgacgggtggcagaccccccgagc	1596
Qy	2044	ggcgcccccccaagagggcggggccacatcacccagccctgcgagtgaggcgctcc	2103
Db	1597	ggcgcccccccaagagggcggggccacatcacccagccctgcgagtgaggcgctcc	1656
Qy	2104	gtcgaacctgagcttctctgtccagcaacaacctgccacctacagagcagctgacccgtg	2163
Db	1657	gtcgaacctgagcttctgtccagcaacaacctgtccacctacagagcagctgacccgtg	1716
Qy	2164	cccaggaggggcccccgatgaggggtccctga	2193
Db	1717	cccaggaggggcccccgatgaggggtccctga	1746

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RESULT 7
US-60-324-185-25008/c
: Sequence 25008, Application US/60324185
: GENERAL INFORMATION:
: APPLICANT: Morris, MacDonald
: APPLICANT: Lal, Preeti
: APPLICANT: Deep, Dinh
: TITLE OF INVENTION: POLYNUCLEOTIDE SEQUENCE DATABASES, AND SINGLE NUCLEOTIDE
: TITLE OF INVENTION: POLYMORPHISMS IDENTIFIED THEREBY
: FILE REFERENCE: GX-0019-1 P
: CURRENT APPLICATION NUMBER: US/60/324,185
: CURRENT FILING DATE: 2001-09-21
: NUMBER OF SEQ ID NOS: 35862
: SOFTWARE: PERL Program
: SEQ ID NO 25008
: LENGTH: 4833
: TYPE: DNA
: ORGANISM: Homo sapiens
: FEATURE:
: NAME/KEY: misc_feature
: OTHER INFORMATION: Incyte ID No: 348127.7
US-60-324-185-25008

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Query Match 51.5%; Score 1637; DB 10; Length 4833;
Best Local Similarity 99.8%; Pred. No. 0;
Matches 1877; Conservative 0; Mismatches 0; Indels 3; Gaps

Qy	549	cttectoatcgctcgtgctgacttccttgactgaacagctgcacccatcgagcagtatgc	608
Db	1883	CTTCCCTCATCGTCCTGCTGTGCCTCATCTTCACGCTGCTGCCACATCGAGCAGTAGC	1824
Qy	609	cgccctggccacgggacacctctcttgagatgagatcgtctggtgggtgtctctcgggac	668
Db	1823	CGCCCTGGCCACCAGGGACACTCTCTTGGAATGGAGATCGCTGTGTGGTTCTTCGGGAC	1764
Qy	569	ggagtagctggttcgcgcctctggttcgcgcgcgcgtgcgcgacgaagtacgtgggcctctggg	728
Db	1763	GGAGTAGCTGCTCGCCTCTGTGTCGCCCGCTGCCGACGAAGTAGTACGTGGGCCCTCTGGGG	1704
Qy	729	gcgcctgcgcctttgcccggaagccaatttcoatacatgaactcatcgttgtcgtggcctc	788
Db	1703	GCGGCTGCGGCTTTTGCCCGGAAGCCCAATTCCAATCATCACCTCAPCTGTGTCGTGGCCCTC	1644
Qy	789	catgggtggctcctgctgctgggtcccagaaggcgaggtgattgscacgtcgcccatcagggg	848
Db	1643	CATGGTGGTCTCTGCGTGGGCTCCAAAGGGGAGGTGTTCACACGCTCGGCCANTCAGGGG	1584
Qy	849	catcgccttcctgcagatccttgagtagtcaacacgtcacgcgcgcagggagggaccctggag	908
Db	1583	CATCCGCTTCTCGACATCCTGAGGATGCTACACGCTCGACCGCCAGGAGGCACCTGGAG	1524
Qy	909	gctccctggctcgtggtcttcacacgcgcgcagagctgataaccaacctgtacatcgg	968
Db	1523	GCCTCTGGGCTCGGTGGTCTTCATCCACGCCAGAGAGTGATAACACCCCTGTACATCGG	1464
Qy	969	cttccctgggcctaatctctcctcgtactttgttacctggctgagaaggagcgcgtgaa	1028
Db	1463	CTWTCTGGGCCTCATCTTCTCTCGTACTTTGTGTACCTTGCTGAGAAGGACGCGGTCAA	1404
Qy	1029	cgaatcaggccgcgtggaatttcggcagctacacagatgcgctgtggtgggggtggtctaac	1088
Db	1403	CGAGTACAGGCCGCGTGGAGTTCCGGAGCTACCGAGTACGCAGATCGCTGTGTGGGGGGTGGTCAC	1344
Qy	1089	agtcaaccaactcggcta tggggacaagggtgccacagctgggtcggggaagaccatacgc	1148
Db	1343	AGTCACCAACCATCGGCTATGSGGACAAGGTGCCCCAGACGTGGGTGGCGGAGACCATCGC	1284
Qy	1149	ctcctgctctctgtcttggcatctctcttcttgctcctccagcgggattcttggctc	1208
Db	1283	CTCCTCGCTTCTGTGCTTTTGGCATCTCCCTCTTTTGGCTCCACGCGGGAATCTTTGGCTC	1224

Qy	2435	ccttggcccccaaatgggtgatgttgacatacacttgcatggttggttggaccagctggcag	2494
Db	403152	ccttggcccccaaatgggtgatgttgacatacacttgcatggttggaccagctggcag	403211
Qy	2495	ggcacaggcctggcccatgtaTggccaggaagttagcacaggctgagtgcaggccaccc	2554
Db	403212	ggcacaggcctggcccatgtaTggccaggaagttagcacaggctgagtgcaggccaccc	403271
Qy	2555	TgctTggccccaggggcttccctgaggggagacagacaacccctggaccccgagctcaaa	2614
Db	403272	TgctTggccccaggggcttccctgaggggagacagacaacccctggaccccgagctcaaa	403331
Qy	2615	Tccaggacctgcacggacagacagcaggcaggacccacccagctgactacagggccaccc	2674
Db	403332	Tccaggacctgcacggacagcaggcaggcaggacccacccagctgactacagggccaccc	403391
Qy	2675	ggcaataaaagccacaggagcccaattTggaggcctgggctgagctccctcactctcagga	2734
Db	403392	ggcaataaaagccacaggagcccaattTggaggcctgggctgagctccctcactctcagga	403451
Qy	2735	aaTgctgaccctaTgggcaggagactTgagagactgctctgagcccccgacttccagcag	2794
Db	403452	aaTgctgaccctaTgggcaggagactTgagagactgctctgagcccccgacttccagcag	403511
Qy	2795	gagggacagctctcaccaattTccccaggggcacgtTggtTgagTgggggggaacgcccccttc	2854
Db	403512	gagggacagctctcaccaattTccccaggggcacgtTggtTgagTgggggggaacgcccccttc	403571
Qy	2855	ctgggtTtagactccagctctctctagctTgagagagagccctTgctctccgccccctgagc	2914
Db	403572	ctgggtTtagactccagctctctctagctTgagagagagccctTgctctccgccccctgagc	403631
Qy	2915	ccactgTgctTgggggtcccgctccaaacccctTgcccagTccagcagccagccaaaca	2974
Db	403632	ccactgTgctTgggggtcccgctccaaacccctTgcccagTccagcagccagccaaaca	403691
Qy	2975	cacagaaggggagctgcacatccccctTgcagctgctgagccgcagagaaTgacggttc	3034
Db	403692	cacagaagggagctgcacatccccctTgcagctgctgagccgcagagaaTgacggttc	403751
Qy	3035	ctacacaggaaggggtctccctctgggcattTacatcgcatagaaaaTcaataattTtgt	3094
Db	403752	ctacacaggaaggggtctccctctgggcattTacatcgcatagaaaaTcaataattTtgt	403811
Qy	3095	gattTgagctgTgttttaagagttTccagTgtgatTttgattTattTattTgtTgcaagc	3154
Db	403812	gattTgagctgTgttttaagagttTccagTgtgatTttgattTattTattTgtTgcaagc	403871
Qy	3155	Ttttcctaataaacgtggagaatcaca	3181
Db	403872	Ttttcctaataaacgtggagaatcaca	403898

RESULT

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US-09-758-466-318
;
; Sequence 318, Application US/09758466
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PM036
; CURRENT APPLICATION NUMBER: US/09/758,466
; CURRENT FILING DATE: 2001-01-11
; PRIOR APPLICATION NUMBER: 60/179,065
; PRIOR FILING DATE: 2000-01-31
; PRIOR APPLICATION NUMBER: 60/180,628
; PRIOR FILING DATE: 2000-02-04
; NUMBER OF SEQ ID NOS: 814
; SOFTWARE: PatentIn ver. 2.0
; SEQ ID NO 318
; LENGTH: 1141
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:

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Qy 1853 accagtcattgggaagccctcactgttctatctcgtctcagaaaaagacaagatcgcg 1912
|||||
Db 901 accagtcattgggaagccctcactgttctatctcgtctcagaaaaagacaagatcgcg 960
|||
Qy 1913 gca 1915
|||
Db 961 gca 963

RESULT 10

US-09-904-809-19624
; Sequence 19624, Application US/09904809
; GENERAL INFORMATION:
; APPLICANT: Hyseq, Inc.
; TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
; FILE REFERENCE: 20411-757
; CURRENT APPLICATION NUMBER: US/09/904, 809
; CURRENT FILING DATE: 2001-07-12
; PRIOR APPLICATION NUMBER: 09/234, 611
; PRIOR FILING DATE: 1999-01-22
; NUMBER OF SEQ ID NOS: 21025
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 19624
; LENGTH: 469
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(469)
; OTHER INFORMATION: n = A,T,C or G
US-09-904-809-19624

Query Match 13.3%; Score 422; DB 7; Length 469;
Best Local Similarity 100.0%; Pred. No. 8.7e-199;
Matches 422; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 2213 gatggcctgagtgagaggagggcgaagtggtggccacacctggccctctctgaaggagg 2272
Db 47 gatggcctgagtgagaggagggcgaagtggtggccacacctggccctctctgaaggagg 106
Qy 2273 ccactctctaaagccagagagagagcccccactctcagagggcccaataccccatgg 2332
Db 107 ccactctctaaagccagagagagagcccccactctcagagggcccaataccccatgg 166
Qy 2333 accatgctgtctggcacagcctgcacttgggggtcagcaaggccacacctctctctggccg 2392
Db 167 accatgctgtctggcacagcctgcacttgggggtcagcaaggccacacctctctctggccg 226
Qy 2393 gtgtggggggcccgctctcaggtctgagttgtttaccccaagcgccctggcccccacatggt 2452
Db 227 gtgtggggggcccgctctcaggtctgagttgtttaccccaagcgccctggcccccacatggt 286
Qy 2453 gatattgacatcactggcatggtggttgggacccagtgagggcagagggcctggccca 2512
Db 287 gatattgacatcactggcatggtggttgggacccagtgagggcagagggcctggccca 346
Qy 2513 tgatggccaggaagtacagagcctgagtcagggccacacctgcttgccagggggct 2572
Db 347 tgatggccaggaagtacagagcctgagtcagggccacacctgcttgccagggggct 406
Qy 2573 tcctgaggggagacagagcaacccctggaccccgacctcaaatccagagccctggcaggc 2632
Db 407 tcctgaggggagacagagcaacccctggaccccgacctcaaatccagagccctggcaggc 466
Qy 2633 ac 2634
Db 467 ac 468

RESULT 11

US-09-909-627-12677

; Sequence 12677, Application US/09909627
; GENERAL INFORMATION:
; APPLICANT: Hyseq, Inc.
; TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
; FILE REFERENCE: 20411-766
; CURRENT APPLICATION NUMBER: US/09/909, 627
; CURRENT FILING DATE: 2001-07-19
; PRIOR APPLICATION NUMBER: 09/277, 227
; PRIOR FILING DATE: 1999-03-23
; NUMBER OF SEQ ID NOS: 23680
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 12677
; LENGTH: 469
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(469)
; OTHER INFORMATION: n = A,T,C or G
US-09-909-627-12677

Query Match 13.3%; Score 422; DB 7; Length 469;
Best Local Similarity 100.0%; Pred. No. 8.7e-199;
Matches 422; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 2213 gatggcctgagtgagaggagggcgaagtggtggccacacctggccctctctgaaggagg 2272
Db 47 gatggcctgagtgagaggagggcgaagtggtggccacacctggccctctctgaaggagg 106
Qy 2273 ccactctctaaagccagagagagagcccccactctcagagggcccaataccccatgg 2332
Db 107 ccactctctaaagccagagagagagcccccactctcagagggcccaataccccatgg 166
Qy 2333 accatgctgtctggcacagcctgcacttgggggtcagcaaggccacacctctctctggccg 2392
Db 167 accatgctgtctggcacagcctgcacttgggggtcagcaaggccacacctctctctggccg 226
Qy 2393 gtgtggggggcccgctctcaggtctgagttgtttaccccaagcgccctggcccccacatggt 2452
Db 227 gtgtggggggcccgctctcaggtctgagttgtttaccccaagcgccctggcccccacatggt 286
Qy 2453 gatgttgacatcactggcatggtggttgggacccagtgagggcagagggcctggccca 2512
Db 287 gatgttgacatcactggcatggtggttgggacccagtgagggcagagggcctggccca 346
Qy 2513 tgatggccaggaagtacagagcctgagtcagggccacacctgcttgccagggggct 2572
Db 347 tgatggccaggaagtacagagcctgagtcagggccacacctgcttgccagggggct 406
Qy 2573 tcctgaggggagacagagcaacccctggaccccgacctcaaatccagagccctggcaggc 2632
Db 407 tcctgaggggagacagagcaacccctggaccccgacctcaaatccagagccctggcaggc 466
Qy 2633 ac 2634
Db 467 ac 468

RESULT 12

US-60-313-371-1497
; Sequence 1497, Application US/60313371
; GENERAL INFORMATION:
; APPLICANT: Ring, Huijun Z.
; APPLICANT: Malsen, Gareth
; APPLICANT: Townley, David
; APPLICANT: Morris, MacDonald
; TITLE OF INVENTION: Single Nucleotide Polymorphisms Associated With ADME Genes
; FILE REFERENCE: GX-0013-5 P
; CURRENT APPLICATION NUMBER: US/60/313, 371
; CURRENT FILING DATE: 2001-08-16
; NUMBER OF SEQ ID NOS: 2447

```

: SOFTWARE: PERL Program
: SEQ ID NO 1497
: LENGTH: 471
: TYPE: DNA
: ORGANISM: Homo sapiens
: FEATURE:
: NAME/KEY: misc_feature
: OTHER INFORMATION: GB:AB015148.1
US-60-313-371-1497

```

Query Match 12.5%; Score 398; DB 10; Length 471;
Best Local Similarity 99.8%; Pred. No. 7.4e-187;
Matches 448; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY	100	tgcgcttcgctgcagctcccggtgcgcgcgcgtgcggcgcccccgcgcagccctcctc	159
Db	1	tcgcttcgctgcagctcccggtgcgcgcgcgtgcggcgcccccgcgcagccctcctc	60
QY	160	gttatggcgcgccctctctcccgccagggcgcgagagaagcgttcggggttcgggcgcgc	219
Db	61	gttatggcgcgccctctctcccgccagggcgcgagagaagcgttcggggttcgggcgcgc	120
QY	220	ctgccagggcccgcgggggcagcgcgcccttgccaaagaatgcacctcttcgctggag	279
Db	121	ctgccagggcccgcgggggcagcgcgcccttgccaaagaatgcacctcttcgctggag	180
QY	280	ctggcgagaggcgcccgcgggcgcgctctacgcgcccctacgcgcgcgcgcgcga	339
Db	181	ctggcgagaggcgcccgcgggcgcgctctacgcgcccctacgcgcgcgcgcgcga	240
QY	340	ggctccgcgcacctcgctcccgcgcgccgcgcgcgcgcgcgcgcgcgcgcgcgcgc	399
Db	241	ggctccgcgcacctcgctcccgcgcgccgcgcgcgcgcgcgcgcgcgcgcgcgcgc	300
QY	400	ggcccgcgccgcgcgtgagccttagaccgcgcgctctccatctaaagcagcgcgccgcg	459
Db	301	ggcccgcgccgcgcgtgagccttagaccgcgcgctctccatctaaagcagcgcgccgcg	360
QY	460	gtgttggcgcgaccacagctccagggcgcgctctaaacttctcgcgcgttccaccggc	519
Db	361	gtgttggcgcgaccacagctccagggcgcgctctaaacttctcgcgcgttccaccggc	420
QY	520	tgaatactcttgattaccacttcgcgct	548
Db	421	tgaatactcttgattaccacttcgcgct	449

RESULT 13
US-09-933-524-85039
; Sequence 85039, Application US/09933524
; GENERAL INFORMATION:
; APPLICANT: Drmanac, Radote T.

APPLICANT: JONES, Lee W.
TITLE OF INVENTION: Novel Nucleic Acid Sequences Obtained
TITLE OF INVENTION: From Various Libraries

FILE REFERENCE: 774
CURRENT APPLICATION NUMBER: US/09/933,524
CURRENT FILING DATE: 2001-08-20
PRIOR APPLICATION NUMBER: 09/528,409
PRIOR FILING DATE: 2000-03-17
NUMBER OF SEQ ID NOS: 116231
SOFTWARE: Hy-patent.pl Version 3.1
SEQ ID NO 85039
LENGTH: 432

```

; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-933-524-85039

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US-09-9333-524-85039

Query Match	10.4%;	Score 332;	DB 9;	Length 432;
Best Local Similarity	100.0%;	Prod. No. 4.6e-154;		
Matches 332;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;
QY 1663	accacatctcacagctgcggaacacacattcggccaccattaaaggttcattcagcagcatg	1722		
Db 101	accacatctcacagctgcggaacacacattcggccaccattaaagtcattcagcagcatg	160		
QY 1723	cagtcactttgtggccaaagagaattccagcaagcgcggaagccttacgattgtcgggagc	1782		
Db 161	cagtcactttgtggccaaagagaattccagcaagcgcggaagccttacgattgtcgggagc	220		
QY 1783	gtcattgaagcagttactcgcaggggcccactcaacctcatgttgcgcatcaaggagctgcag	1842		
Db 221	gtcattgagcagttactcgcaggggcccactcaacctcatgttgcgcatcaaggagctgcag	280		
QY 1843	aggaggttggaaccagttccattgggaagccctcactgttcattctccgtctcagaaaaagc	1902		
Db 281	aggaggttggaaccagttccattgggaagccctcactgttcattctccgtctcagaaaaagc	340		
QY 1903	aagatctcggcgacgacacacagatcggcgcccgctgaaccgagttagaagacaaggtgcag	1962		
Db 341	aagatctcggcgacgacacacagatcggcgcccgctgaaccgagttagaagacaaggtgcag	400		
QY 1963	cagctgacacagaaggctgggaactcatcacgga	1994		
Db 401	cagctgacacagaaggctgggaactcatcacgga	432		

RESULT : 14

US-09-933-524-18451

; Sequence 18451, Appl

GENERAL INFORMATION:
APPLICANT: Drmanac

APPLICANT: Labat, Ivan

; APPLICANT: Stache-Crain, Birgit

; APPLICANT: Dickson, Mark

APPLICANT: JONES, Lee W.
TITLE OF INVENTION: Novel

; TITLE OF INVENTION: From Various Libraries

; FILE REFERENCE: 774

; CURRENT APPLICATION NUMBER: US/09/933,524
 ; CURRENT FILING DATE: 2001-08-30

; PRIOR APPLICATION NUMBER: 09/528,409

; PRIOR FILING DATE: 2000-03-17

NUMBER OF SEQ ID NOS: 116231

: SOFWAKE: HY-P
: SEO ID NO 18451

; LENGTH: 455

; TYPE: DNA

; ORGANISM: Homo sapiens
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Query Match	9.9%	Score 316;	DB 9;	Length 455;
Best Local Similarity	99.7%;	Pred. No. 4,1e-146;		
Matches 366;	Conservative	0;	Mismatches 1;	Indels 0;
				Gaps 0;

Qy 1817 tcatggtcgcacatcaaggagctcacaqaggaqgcttqaccagtccatttqqqaagccctcac 1876

Db 69 tcatggtgcgcatacaggagctgcagaggaggctggaccagtcattgggaagccctcac 128

Qy 1877 tqtccatctccqtctcagaaaaagacaaqqatccqccacacacacqatccqccccc 1936

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Db 129 tggtcatctccgtctctcagaaaagagcaaggatcgcggcagcaaacacgacgacgccc 188

Qy 1937 tgaaccgaatagaagacaaaggtgacgcagctggaccgaaggtggcactcatcaccgaca 1996

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Db 189 tgaccgagtagaagacaagtgacgcacctggaccagaggctggcactcatcaccgaca 248

Qy 1997 tacttaccacactactctccctgtcacacctgccacaccccccccagggcgcgccccca 2056

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Db 249 tgcttcaccagctgctctctcttgcacggtggcagcaccctccggcagcggcgccccca 308

Qy 2057 gagaggcgcccccacacacccagccctggcagtgggcgtccgtcgacccctgagc 2116
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Qy 309 gagaggcgcccccacacacccagccctggcagtgggcgtccgtcgacccctgagc 368
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Qy 2117 tcttctgcccagcaacacccctgcccacacacagcagcagcgtgcccagagggcc 2176
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Qy 369 tcttctgcccagcaacacccctgcccacacacagcagcagcgtgcccagagggcc 428
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Qy 2177 ccgatga 2183
Db |||||||
Qy 429 ccgatga 435
Db |||||||

RESULT 15
US-09-904-703-5165
; Sequence 5165, Application US/09904703
; GENERAL INFORMATION:
; APPLICANT: Hyseq, Inc.
; TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
; FILE REFERENCE: 20411-758CON1
; CURRENT APPLICATION NUMBER: US/09/904,703
; CURRENT FILING DATE: 2001-07-12
; PRIOR APPLICATION NUMBER: 09/210,298
; PRIOR FILING DATE: 1998-12-09
; NUMBER OF SEQ ID NOS: 17812
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 5165
; LENGTH: 412
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-904-703-5165

Query Match 9.7%; Score 307; DB 7; Length 412;
Best Local Similarity 99.5%; Pred No. 1.2e-141;
Matches 407; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy 2421 tgttaccacaaagcgcctggccccacatggtgatgttgacatcactggcatggtggtg 2480
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Qy 2481 ggaccagtgccagggcacagggcctggcccatgtatggccaggaagtagcacaggtga 2540
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Qy 64 ggaccagtgccagggcacagggcctggcccatgtatggccaggaagtagcacaggtga 123
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Qy 2541 gtgagggccacccctgcttgcccaggggcttctctgagggagacagagcaacccctgg 2600
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Qy 124 gtgagggccacccctgcttgcccaggggcttctctgagggagacagagcaacccctgg 183
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Qy 2601 acccagcctcaaatccaggacccctggccagggcacagggcagggcagccacagcgtg 2660
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Qy 184 acccagcctcaaatccaggacccctggccagggcacagggcagggcagccacagcgtg 243
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Qy 2661 actacagggccacccggcaataaaagccacagggccatttgagggcctggcctggctc 2720
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Qy 244 actacagggcccgccggcaataaaagccacagggccatttgagggcctggcctggctc 303
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Qy 2721 cctcactctcaggaatgctgacctgagggcagggagactgtgagagactgctcctgagcc 2780
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Qy 304 cctcactctcaggaatgctgacctgagggcagggagactgtgagagactgctcctgagcc 363
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Qy 2781 ccagcttccagcagggagcaggtctccatttccccagggcagcgtg 2829
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
Qy 364 ccagcttccagcagggagcaggtctccatttccccagggcagcgtg 412
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||||||

Search completed: November 2, 2001, 15:05:20
Job time: 6746 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: November 2, 2001, 12:09:44 ; Search time 1660.94 Seconds
(without alignments)
18103.921 Million cell updates/sec

Title: US-09-135-010A-1
Perfect score: 3181
Sequence: 1 ctgccctctcggcccccgc.....aataaacgtggagaatcaca 3181

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 10228115 seqs, 4726426750 residues

Word size : 12

Total number of hits satisfying chosen parameters: 2312605

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database :

EST:*

1: gb_est1:*
2: gb_est2:*
3: gb_est3:*
4: gb_est4:*
5: gb_est5:*
6: gb_est6:*
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181: em_estom3: *
182: em_estpl11: *
183: em_estro21: *
184: em_estro22: *
185: em_estro23: *

190: gb_estl10: *
191: gb_estl11: *
192: gb_htc: *
193: em_gss_fun: *
194: em_gss_hum1: *
195: em_gss_hum2: *
196: em_gss_hum3: *
197: em_gss_hum4: *
198: em_gss_hum5: *
199: em_gss_hum6: *
200: em_gss_hum7: *
201: em_gss_hum8: *
202: em_gss_hum9: *
203: em_gss_inv1: *
204: em_gss_inv2: *
205: em_gss_inv3: *
206: em_gss_other: *
207: em_gss_pln1: *
208: em_gss_pln2: *
209: em_gss_pro: *
210: em_gss_rod1: *
211: em_gss_rod2: *
212: em_gss_rod3: *
213: em_gss_rod4: *
214: em_gss_rod5: *
215: em_gss_vrt1: *
216: em_gss_vrt2: *
217: em_gss_vrt3: *
218: gb_gss1: *
219: gb_gss2: *
220: gb_gss3: *
221: gb_gss4: *
222: gb_gss5: *
223: gb_gss6: *
224: gb_gss7: *
225: gb_gss8: *
226: gb_gss9: *
227: gb_gss10: *
228: gb_gss11: *
229: gb_gss12: *
230: gb_gss13: *
231: gb_gss14: *
232: gb_gss15: *
233: gb_gss16: *
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247: gb_gss30: *
248: gb_gss31: *
249: gb_gss32: *
250: gb_gss33: *
251: gb_gss34: *
252: em_gss_inv4: *
253: em_gss_rod6: *
254: em_gss_rod7: *
255: em_gss_rod8: *
256: gb_gss35: *
257: gb_gss36: *
258: gb_gss37: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

Genoscope - Centre National de Sequencage
BP 191 91006 EVRY cedex - France
Email: seqref@genoscope.cns.fr, Web : www.genoscope.cns.fr.
Location/Qualifiers

FEATURES

source 1. 851
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="CS0DK007YA04"
/tissue_type="placenta"
/note="Vector: pCMVSPORT 6; Site: 1: NotI; 1st strand cDNA was primed with a NotI-oligo(dT) primer. Five prime end enriched, double-stranded cDNA was digested with NotI and cloned into the NotI and EcoRV sites of the pCMVSPORT 6 vector. Library was normalized. Library was constructed by Life Technologies. Contact : Feng Liang Life Technologies, a Division of Invitrogen 9800 Medical Center Drive, Rockville, Maryland 20850, USA Fax : (1) 301 610 8371
Email : fliang@lifetech.com URL :
http://fulllength.invitrogen.com"

BASE COUNT 154 a 249 c 273 g 173 t 2 others
ORIGIN

Query Match 22.9%; Score 727; DB 106; Length 851;
Best Local Similarity 99.8%; Pred. No. 0;
Matches 827; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2289 ccagagaaagagcccccactctcagagggcccaataccccatggaccatgctctggca 2348
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Db 841 CCAGAGAGAAGAGCCCACTCTCAGAGGCCCAATACCCATGGACCATGCTGTGGMA 782
QY 2349 cagcctgcacttggggctcagagggccacactcttctcggcggtgtggggcccccgc 2408
|||||
Db 781 CAGCCTGCTGCTGGGGCTCAGACAGGCCACCTCTTCTTGGCGGTGGGGCCCGGTC 722
QY 2409 tcagctcgtgagttgtaccccaagcgccctggccccacatcgtgtagttgacatcactg 2468
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Db 721 TCAGGCTCTGAGTGTGTACCCCAAGCGCCCTGGCCCCACATGGTGATGTTGACATCAGTG 662
QY 2469 gcattggttggggcccaagtcaggggcaagggcctggccctatgattgcccagaagt 2528
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Db 661 GCATGGTGTGGGACCCAGTGGGACAGGCGCTGGCCCATGATATGGCCAGGAAGT 602
QY 2529 agcacagctcagtcagggccacactcctgtcggccaggggcttctcaggggagacag 2588
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Db 601 AGCAGAGCTGAGTGCAGGCCACCTGCTTGGCCAGGGGGCTTCTTGGGGAGACAG 542
QY 2589 agcaacccttgagcccccagcctcaaatccagagaccctgcagggcacaggcagggac 2648
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Db 541 AGCAACCCCTGGACCCAGCCTCAATCCAGGACCCCTGCCAGGCACAGCGGAGGAC 482
QY 2649 cagccacgctgactacagggccacgggcaataaaagccagagccatttggagggcc 2708
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Db 481 CAGCCACGCTGACTACAGGGCGCGGCAATAAAAGCCAGAGGCCCATTTGGAGGGCC 422
QY 2709 tgggcttggtccctcactctcaggaaatgctgacccatggcagggagagactgagagact 2768
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Db 421 TGGGCTGGCTCCCTCACTCTCAGAAATGCTGACCCATGGCAGGAGACTGTGGAGACT 362
QY 2769 gctcctgagcccccagcttccagggagggagcagctctcaccatttcccccagggcagctg 2828
|||||
Db 361 GCTCCTGAGCCCCCAGCTTCCAGCAGGAGGAGGAGTCTCACCATTTCACAGGGGACGGTG 302
QY 2829 gttgagtggggggaaagcccccacttccctgggttagactgccagcttcttccagctgaga 2888
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Db 301 GTTGTAGTGGGGGAAAGCCCACTTCCCTGGTGTAGACTGCCAGCTTCTTCTAGCTGAGCA 242
QY 2889 ggaagccttgcctctccccccttgagcccaacttgagtggtggggtcccgctcccaacccctc 2948
|||||
Db 241 GGAGCCCTGCCTCTCTCGGCCCTGAGCCCACTGTGGGTGGGGCTCCCGCCCTCCCAACCCCTC 182
QY 2949 gccagtcgccagccagccagcaaacacacagaagggggactgccacctcccccttgcagct 3008

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
C 1	727	22.9	851	106	AL577771	AL577771 AL577771
C 2	529	16.6	679	138	BE675840	BE675840 7f17a09.x
C 3	514	16.2	600	108	AU141948	AU141948 AU141948
C 4	511	16.1	943	106	AL577772	AL577772 AL577772
C 5	507	15.9	812	152	BG328061	BG328061 602427108
C 6	497	15.6	775	153	BG385872	BG385872 602454417
C 7	491	15.4	491	110	AW006385	AW006385 wt04g12.x
C 8	474	14.9	488	113	AW205864	AW205864 UI-H-B11-
C 9	461	14.5	461	159	AI1347525	AI1347525 q980612.x
C 10	442	13.9	442	24	AI1769029	AI1769029 wg31h01.x
C 11	441	13.9	494	112	AW192638	AW192638 x14b04.x
C 12	440	13.8	493	20	AI439544	AI439544 tc90b05.x
C 13	408	12.8	484	110	AW005483	AW005483 ws94f02.x
C 14	393	12.4	451	144	BF109081	BF109081 7150b06.x
C 15	387	12.2	418	19	AI344361	AI344361 qp07e12.x
C 16	378	11.9	684	155	BG548631	BG548631 602576435
C 17	367	11.5	535	169	BF798694	BF798694 RC1-C1011
C 18	363	11.4	481	24	AI738525	AI738525 w133a11.x
C 19	363	11.4	556	108	AU159371	AU159371 AU159371
C 20	346	10.9	683	19	AI344314	AI344314 tc03e03.x
C 21	333	10.5	631	164	BE159001	BE159001 MR0-HT040
C 22	318	10.0	490	24	AI739442	AI739442 w113e04.x
C 23	313	9.8	328	102	AI826701	AI826701 wk46e04.x
C 24	305	9.6	318	112	AW136422	AW136422 UI-H-B11-
C 25	269	8.5	371	121	AW866858	AW866858 CM3-SN003
C 26	255	8.0	460	165	BE222815	BE222815 hu53g08.x
C 27	253	8.0	253	19	AI344116	AI344116 tc02b06.x
C 28	251	7.9	339	9	AA603649	AA603649 np20g12.s
C 29	251	7.9	476	12	AA824263	AA824263 a129e05.s
C 30	242	7.6	565	191	W93500	W93500 zd96e05.s1
C 31	233	7.3	244	169	BF755715	BF755715 PM4-CT056
C 32	232	7.3	233	111	AW057846	AW057846 wv90a10.x
C 33	213	6.7	377	103	AI869254	AI869254 w15e10.x
C 34	206	6.5	323	19	AI344946	AI344946 tb01c01.x
C 35	206	6.5	437	19	AI344925	AI344925 tb01a01.x
C 36	206	6.5	442	19	AI344927	AI344927 tb01a03.x
C 37	205	6.4	258	19	AI345107	AI345107 t895e07.x
C 38	205	6.4	391	20	AI473902	AI473902 ti68c11.x
C 39	205	6.4	1003	138	BE617784	BE617784 601441990
C 40	202	6.4	449	122	AW953589	AW953589 EST365659
C 41	200	6.3	405	123	AW978648	AW978648 EST3909757
C 42	197	6.2	197	112	AW138991	AW138991 UI-H-B11-
C 43	196	6.2	309	6	AA352245	AA352245 EST60369
C 44	196	6.2	465	122	AW949927	AW949927 EST361997
C 45	184	5.8	270	113	AW268275	AW268275 xr95a03.x

ALIGNMENTS

RESULT 1
AL577771/c
LOCUS AL577771 LTI_NFL006_PL2 Homo sapiens cDNA clone CS0DK007YA04 3
DEFINITION prime, mRNA sequence.
ACCESSION AL577771
VERSION AL577771.1 GI:12941220
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 851)
AUTHORS Li, W.B., Gruber, C., Jesse, J. and Polayes, D.
TITLE Full-length cDNA libraries and normalization
JOURNAL Unpublished (2001)
COMMENT Contact: Genoscope


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Db 181 GCCAGTCCAGCAGCCAGCAACACACAGAGGGGACTGCCACCTCCCTTGCCAGCT 122
Qy 3009 gctgagccgcagagagtgacggttctctacacagaggggtctctctggtgacattaca 3068
Db 121 GCTGAGCCGCAGAGAGTGCAGGTTCTTACACAGACAGAGGGTTCTCTGTGGCATTACA 62
Qy 3069 tcgcataaatacaataatttggtgattggatctggttttaataga 3117
Db 61 TCGCATAGAAATCAATATTTGGTGATTTGGATCTGTGTTTAATGA 13

RESULT 2
BE675840 679 bp mRNA EST 08-SEP-2000
LOCUS 7f17a09.x1 NCI-CGAP-CLL1 Homo sapiens cDNA clone IMAGE:3294904 3'
DEFINITION similar to TR:060607 O60607 SLOW DELAYED RECTIFIER CHANNEL SUBUNIT.
; contains PTR5.L3 TAR1 repetitive element ; mRNA sequence.
ACCESSION BE675840
VERSION BE675840.1 GI:10036381
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 679)
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
JOURNAL Contact: Robert Strausberg, Ph.D.
COMMENT Email: cgapb-r@mail.nih.gov
Tissue Procurement: Ash Alizadeh, John Byrd, M.D., Mike Grever,
M.D., Louis M. Staudt, M.D., Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D.
DNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL, send email to:
info@image.llnl.gov
Seq primer: -40UP from Gibco
High quality sequence stop: 465.
FEATURES
source
Location/Qualifiers
1..679
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:3294904"
/clone_lib="NCI-CGAP-CLL1"
/tissue_type="B-cell, chronic lymphocytic leukemia"
/lab_host="DH10B"
/notes="vector: p7T3D-Pac (Pharmacia) with a modified
polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA
was primed with a Not I - oligo(dT) primer [5'
TGTTACCAACTGAAGTGGGCGCGCGCATGCTTTTCTTTTCTTTTCTTTTCTTTT
T 3']; double-stranded cDNA was ligated to Eco RI
adaptors (Pharmacia), digested with Not I and cloned into
the Not I and Eco RI sites of the modified p7T3 vector.
Library is normalized, and was constructed by Bento
Soares and M.Fatima Bonaldo."
BASE COUNT 93 a 224 c 207 g 155 t
ORIGIN

Query Match 16.6%; Score 529; DB 138; Length 679;
Best Local Similarity 99.7%; Pred. No. 2.2e-254;
Matches 629; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy 405 gcggccgcggtagcctagaccgcgcgctctctctacacagcgcgcgcggtgtt 464
Db 1 CGCGCCGCCGTGAGCCCTAGACCCGCGCTCTCCATCTACAGCAGCGCGCGCGGTGT 60
Qy 465 ggcgcgaccacgctccagggccgcgctctctctctctctctctctctctctctct 524
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Db 61 GGCGCGCACCCACGTCACGAGGCGCGGTCACAACTTCTCGAGCGTCCACCGGCTGAA 120
Qy 525 atgcttggtttacacacttcgcccgtctctctctctctctctctctctctctctct 584
Db 121 ATGCTTCGTTTACACATTCGCCGCTTTCCTTCATCGTCTGCTGCTGCTGCTGCTG 180
Qy 585 gctgctccaccatcgacagcagtatccgcccctggccacggggaactctctctctct 644
Db 181 GCCTGTCACCATCGAGCAGTATGCCGCCCTGGCCACGGGACTCTCTCTGGATGGAG 240
Qy 645 cgtgctgggtggttctctctgggacggagtagctggtgcgcctctggtccgcggctcc 704
Db 241 CGTGTGGTGGTGTCTTTCGAGGAGTACGTGGTCCGCTCTGGTCCGCGGCTGCCG 300
Qy 705 cagcaagtcagtggtggtctctggggggtggtggtggtggtggtggtggtggtggt 764
Db 301 CAGCAAGTACGTGGGCTCTGCGGCGGCTTTCCTTCCTTCCTTCCTTCCTTCCTTC 360
Qy 765 cgacctcatgctggtggtggtggtggtggtggtggtggtggtggtggtggtggtg 824
Db 361 CGACCTCATCGTGGTGTGTCGCTCCATGCTGCTGCTGCTGCTGCTGCTGCTGCTG 420
Qy 825 gttgtccagtcggtggtggtggtggtggtggtggtggtggtggtggtggtggtggt 884
Db 421 GTTGTGCACGTGCGCCATCAGGGGCTTCCTTCCTTCCTTCCTTCCTTCCTTCCT 480
Qy 885 cgaccgccaggggacgtggtggtggtggtggtggtggtggtggtggtggtggtggtg 944
Db 481 CGACCGCCATCGAGGACCTGAGGCTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 540
Qy 945 gctgataaccacccctgatacgtggtggtggtggtggtggtggtggtggtggtggt 1004
Db 541 GCTGATTAACACACCTGATCATCGCTTCTTCCTGGGCTCATCTCTCTCTCTCTCT 600
Qy 1005 cctggtcgagaagcgcggtgacagtcga 1035
Db 601 CTGTGCTGAGAGGACGCGGTGAGACGAGTCA 631

RESULT 3
AUI141948
LOCUS AUI141948 600 bp mRNA EST 25-OCT-2000
DEFINITION AUI141948 THYROI Homo sapiens cDNA clone THYROI001510 5', mRNA
sequence.
ACCESSION AUI141948
VERSION AUI141948.1 GI:11003469
KEYWORDS EST.
SOURCE Homo sapiens
ORGANISM human.
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 600)
AUTHORS Ota, T., Nishikawa, T., Suzuki, Y., Ishii, S., Saito, K., Kawai, Y.,
Yamamoto, J., Wakamatsu, A., Nakamura, Y., Nagai, T., Sugano, S. and
Isogai, T.
TITLE HRI human cDNA project
JOURNAL HRI human cDNA project; 5'- & 3'-end one pass sequencing: Helix
COMMENT Unpublished (2000)
Contact: Takao Isogai
Genomics Laboratory
Helix Research Institute
1532-3 Yana, Kisarazu, Chiba 252-0812, Japan
Tel: 81-438-52-3951
Fax: 81-438-52-3952
Email: genomics@hri.co.jp
HRI human cDNA project; 5'- & 3'-end one pass sequencing: Helix
Research Institute; cDNA library construction: Department of
Virology, Institute of Medical Science, University of Tokyo, and
Helix Research Institute.
FEATURES
Location/Qualifiers
1..600
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="THYROI001510"
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FEATURES

REFERENCE 1 (bases 1 to 812)
NIH-MGC http://mgc.nci.nih.gov/
AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)
TITLE Unpublished (1999)
JOURNAL
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: Ling Hong/Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: NIH Intramural Sequencing Center
VERSION Clone distribution: MGC clone distribution information can be
KEYWORDS found through the I.M.A.G.E. Consortium/LLNL at:
SOURCE http://image.llnl.gov
ORGANISM Plate: LLC1233 row: o column: 02
High quality sequence stop: 725.
FEATURES Location/Qualifiers
source 1..812
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:454677"
/clone_lib="NIH_MGC_15"
/tissue_type="adenocarcinoma cell line"
/lab_host="DH10B (phage-resistant)"
/note="Organ: colon; Vector: pORF7; Site_1: xhoI; Site_2: EcoRI; CDNA made by oligo-dt priming. Directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GGCACGAG(G). Size-selected >500bp for average insert size 1.8kb. Library constructed by Ling Hong in the laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies)"
BASE COUNT 193 a 234 c 233 g 152 t
ORIGIN
Query Match 15.9%; Score 507; DB 152; Length 812;
Best Local Similarity 100.0%; Pred. No. 2.6e-243; Mismatches 0; Indels 0; Gaps 0;
Matches 507; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 998 ttgttacctggtgaggaagcgggtgaacgagtcagcgccggtgaggttcgagct 1057
Db 18 TTGTCTACTTGGCTGAGAAGGACGCGGTGAAGAGTCAGCGCGGTGGAGTTCGCGAGT 77
Qy 1058 acgagatgcgctgtggtggtggtggtggtggtggtggtggtggtggtggt 1117
Db 78 ACGAGATGCGTGTGTGGGGGTGGTGCACAGTCACACCATCGCTATGGGACAAAG 137
Qy 1118 tgccccagactggttcgggaagaccatcgctctgcttctctgttgcattctct 1177
Db 138 TGCCCCAGACGTGGGTGCGGAAGACCATCGCTCTCTGTCTGTCTGTCTCTCT 197
Qy 1178 tctttgcctccagcggggtattctgtcggtggttgcctgaagtgagcagagaagc 1237
Db 198 TCTTTGGCTCCACGCGGGATCTTGTGCTCGGGGTTCCTTGTGCTGAGGTGACGAGAAGC 257
Qy 1238 agaggcagaagcaattcaacccggcagatcccggtggtggtggtggtggtggtggt 1297
Db 258 AGAGCGAGAAGCACTTAACCGGAGATCCCGGGCGGAGCTCACTTCACTTCAAGCCGAT 317
Qy 1298 ggaagtgtatgtgcgagaaaccccgactcctcaactggaagtctatattccggaagg 1357
Db 318 GGAGGTGCTATGCTGCGGAGAACCCCGACTCTCCACCTGGAAGATCTACATCCGGAAG 377
Qy 1358 cccccggagccacactctgcttaccagcccccaaaccaagactggtggtggtggtggt 1417
Db 378 CCCCCGGAGCCACACTCTGCTGTACCCAGCCCCCAACCAAGAGTGTGTGGGTAA 437
Qy 1418 agaaaaaaagtccaagctgacaaagacaatgggtgactctctgagagaagatgctca 1477
Db 438 AGAAAAAAGTTCAAGCTGGACAAAGACAATGGSGTGACTCTCTGGAGAGAGATGCTCA 497
Qy 1478 cagtcctccatatacagtcgaccccc 1504
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Db 498 CAGTCCCCCATATACAGTGGCAGCCCC 524
RESULT 6
BG385872
LOCUS BG385872
DEFINITION 775 bp mRNA EST 12-MAR-2001
602454417F1 NIH_MGC_15 Homo sapiens cDNA clone IMAGE:4582565 5',
mRNA sequence.
ACCESSION BG385872
VERSION BG385872
KEYWORDS GI:13279276
SOURCE EST.
ORGANISM human.
REFERENCE Homo sapiens
1 (bases 1 to 775)
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
TITLE NIH-MGC http://mgc.nci.nih.gov/
JOURNAL National Institutes of Health, Mammalian Gene Collection (MGC)
COMMENT Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: Ling Hong/Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: NIH Intramural Sequencing Center
VERSION Clone distribution: MGC clone distribution information can be
KEYWORDS found through the I.M.A.G.E. Consortium/LLNL at:
SOURCE http://image.llnl.gov
ORGANISM Plate: LLC1305 row: b column: 06
High quality sequence stop: 772.
FEATURES Location/Qualifiers
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/db_xref="taxon:9606"
/clone="IMAGE:4582565"
/clone_lib="NIH_MGC_15"
/tissue_type="adenocarcinoma cell line"
/lab_host="DH10B (phage-resistant)"
/note="Organ: colon; Vector: pORF7; Site_1: xhoI; Site_2: EcoRI; CDNA made by oligo-dt priming. Directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GGCACGAG(G). Size-selected >500bp for average insert size 1.8kb. Library constructed by Ling Hong in the laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies)"
BASE COUNT 157 a 258 c 237 g 123 t
ORIGIN
Query Match 15.6%; Score 497; DB 153; Length 775;
Best Local Similarity 99.8%; Pred. No. 2.7e-238;
Matches 617; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
Qy 2224 gtgagaggaggagcgaagtggtggtggtggtggtggtggtggtggtggtggt 2283
Db 2 GTGAGAGGGAGGCCAAGAGTGGCCCACTGCGCCCTCTCTGAAGGAGGCCACTCTCTAA 61
Qy 2284 aaggccccagagagaagagccactctcagagagccccataccccatgaccatgctgc 2343
Db 62 AAGGCCAGAGAGAGAGGCCCACTCTCAGAGCCCCCAATACCCCATGGACCATGTCTC 121
Qy 2344 tgccacagctcacttggtggtggtggtggtggtggtggtggtggtggtggtggt 2403
Db 122 TGGCAGACCTTGACTTGGGGGCTCAGAGGCCACTCTCTCTGCTGCTGCTGCTGCTG 181
Qy 2404 ccgtctcaggtctgagttgttaccacagcgccttggtggtggtggtggtggtggtggt 2463
Db 182 CCGTCTCAGTGTGAGTTGTTACCCCAAGGCCCTTGCCCCACATGGTGTGATGTGACAT 241
Qy 2464 cactggcattggtggtggtggtggtggtggtggtggtggtggtggtggtggtggtggt 2523
Db 242 CACTGGCATGGTGGTGGGACCCAGTGGGAGGACAGGCGCTGGCCCATGTATGGCCAG 301

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QY 2524 gaagtagacaggtgagtgacagccacccctggtggccagggggtcttctgagggga 2583
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Db 302 GAAGTAGACAGGCTGAGTGACAGCCACCTGCTGGCCCA-GGGGCTTCTGAGGGGA 360
|||||
QY 2584 gacagagcaacccctgacccagcctcaaatcagagccctgcagagcacagcagggc 2643
|||||
Db 361 CACAGACAACCCCTGGACCCACCCCTCAATCCAGAGACCTGCCAGCAGCAGCAGGGC 420
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QY 2644 agaccagccacagctgactacagggccacccggccaataaaagccagagccatttggga 2703
|||||
Db 421 AGGACACGCCACAGCTGACTACAGGGCCACCGGCAATAAAAGCCAGGAGCCCATTTGGA 480
|||||
QY 2704 gggcctgggctggctccctcaactctcaggaatactgacccatgggcagagactgtgg 2763
|||||
Db 481 GGGCCTGGGCTGGCTCCCTCACTCTCAGGAATGCTGACCCATGGGCGAGGAGACTGTGG 540
|||||
QY 2764 agactgctctgagcccccagcttcacagcagagggacagtctcaccatttccccagggc 2823
|||||
Db 541 AGACTGCTCTGACCCCCCAGCTTCCAGCAGGAGGAGAGTCTCACCATTTCGCCAGGGC 600
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QY 2824 acgtgggtgagtgaggggg 2841
|||||
Db 601 ACGTGGTGTAGTGGGGG 618
|||||

RESULT 7
AW006385/c 491 bp mRNA EST 08-MAR-2000
LOCUS wt04g12.x1 NCI_CGAP_C03 Homo sapiens cDNA clone IMAGE:2506534 3'
DEFINITION similar to contains PTR5.b2 TAR1 repetitive element ;, mRNA
sequence.
ACCESSION AW006385
VERSION AW006385.1 GI:5855163
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 491)
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgabbs-r@mail.nih.gov
Tissue Procurement: Elias Campo, M.D., Michael R. Emmert-Buck, M.D.
, Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D.
cDNA Library Arraying: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
Insert length: 686 Std Error: 0.00
Seq primer: -400P from Gibco
High quality sequence stop: 455.
Location/Qualifiers
1. .491
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2506534"
/clone_lib="NCI_CGAP_C03"
/sex="pooled"
/tissue_type="colon"
/lab_host="DH10B"
/notc="Vector: pT73D-Pac (Pharmacia) with a modified
polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA
was prepared from 12 pooled bulk tumor samples and primed
with a Not I - oligo(dT) primer. Double-stranded cDNA was
ligated to Eco RI adaptors (Pharmacia), digested with Not
I and cloned into the Not I and Eco RI sites of the
modified pT73 vector. Library went through one round of
```

```
normalization.
BASE COUNT 111 a 133 c 145 g 102 t
ORIGIN
Query Match 15.4%; Score 491; DB 110; Length 491;
Best Local Similarity 100.0%; Pred. No. 2.7e-235;
Matches 491; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2690 gagccccatttgagggcctggcctcctcactctcaggaatactgaccatgg 2749
|||||
Db 491 GGAGCCCATTTGGAGGGCTGGCGCTGGCTCCTCACTCTCAGGAAATGCTGACCCATG 432
|||||
QY 2750 gcagagagactgtggagactgtcctcagccccagcagcttcacagcagaggagctcac 2809
|||||
Db 431 GCAGGAGAGACTGTGGAGACTGCTCCTCAGCCCCCAGCTTCCAGCAGGAGGAGACTCTCAC 372
|||||
QY 2810 catttccccagggcagctgggttgagtgagggaagcccaacttccctgggttagactgc 2869
|||||
Db 371 CATTTTCCCCAGGGCAGCTGGTTGAGTGGGGGGAAGCCCCACCTCCCTGGGTTAGACTGCC 312
|||||
QY 2870 agctcttctcagtgagagagcctcctcctcgcgcctcagccctcagccactgtcgtgggg 2929
|||||
Db 311 AGCTCTTCTAGCTGAGAGAGAGCCCTGCCTCTCCGCCCTCAGCCCACTGTGCGTGGGG 252
|||||
QY 2930 ctccgcctccaaacccctcgcagtcctccagcagccagccaaacacacagaggggactg 2989
|||||
Db 251 CTCGCCCTCCAAACCCCTCGCCAGTCCCGCAGCAGCCAGCCAAACACACAGAGGGGACTG 192
|||||
QY 2990 ccaactccccctggcagctgtgagccgcagagagagtgacgttctctacacagagcaggg 3049
|||||
Db 191 CCACCTCCCTTCCAGCTGCTGAGCCCGCAGAGAGTGACGGTTCTTACACAGGACAGGG 132
|||||
QY 3050 gtctcttctgggcatcacatcagatcaataattgttggtgattgtgactctgt 3109
|||||
Db 131 GTTCCTCTGGGCATTACATCCATAGAAATCAATAATTTGTGGTGATTTGGATCTGTGT 72
|||||
QY 3110 ttaatgagttcagcagtgattttgattattattgtgcagcttttccataataaacg 3169
|||||
Db 71 TTTAATGAGTTTACAGTGTGATTTTGTATTATTAATTTGCAAGCTTTTCTTAATAAAGC 12
|||||
QY 3170 tgagagaatacac 3180
|||||
Db 11 TGGAGAAATCAC 1
|||||

RESULT 8
AW205864/c 488 bp mRNA EST 02-DEC-1999
LOCUS UI-H-B11-afv-c-12-0-UI-s1 NCI_CGAP_Sub3 Homo sapiens cDNA clone
DEFINITION IMAGE:272967 3', mRNA sequence.
ACCESSION AW205864
VERSION AW205864.1 GI:6505268
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 488)
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgabbs-r@mail.nih.gov
The sequence contained an oligo-dT track that was present in the
oligonucleotide that was used to prime the synthesis of first
strand cDNA and therefore this may represent a bonafide poly A
tail. cDNA Library Preparation: M.B. Soares Lab Clone distribution:
NCI-CGAP clone distribution information can be found through the
I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
Seq primer: M13 Forward
```



```
source          1. 494
                /organism="Homo sapiens"
                /db_xref="taxon:9606"
                /clone="IMAGE:2677903"
                /clone_lib="NCI_CGAP_Pan1"
                /tissue_type="adenocarcinoma"
                /lab_host="DH10B"
                /note="Organ: pancreas; Vector: pCMV-SPORT6; Site_1: SalI;
                Site_2: NotI; Cloned unidirectionally. Primer: Oligo dT.
                Average insert size 1.72 kb. Life Technologies catalog #:
                11548-013"
BASE COUNT      111 a 133 c 144 g 105 t      1 others
ORIGIN

Query Match      13.9%; Score 441; DB 112; Length 494;
Best Local Similarity 99.8%; Pred. No. 3.7e-210;
Matches 491; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2690 ggagccatttgaggcgctggcgctgctccctcactcaggaatgctgacccatgg 2749
Db 494 GGAGCCATTGGAGGGCGCTGGCGCTGCTCCTCCTCCTCAGGAATGCTGACCCATGG 435
QY 2750 gcaggagactgagagactgctctgagcccccagcttccagcagaggagagctctcac 2809
Db 434 GCAGGAGACTGTGGAGACTGCTCTGAGCGCCCGCCAGCTTCCAGCAGGAGGGACAGTCTCAC 375
QY 2810 catttcccagggcacgtggttgagtggggggaaacgcccacttccctggttagactgcc 2869
Db 374 CATTTCCNACGGGCACGTTGTTGAGTGGGGGGAAGCGCCACTTCCCTGGTTAGACTGCC 315
QY 2870 agctcttctagctggagagagccctgcctctccgcccctgagccactgtcgtgggg 2929
Db 314 AGCTCTTCTAGCTGGAGAGAGCGCTGCCCTCTCGGCCCTGAGCCCACTGTGCGTGGGG 255
QY 2930 ctccgctctcaacccctcgccagctccagcagccagccagccaaacacagaggagactg 2989
Db 254 CTCGGCGCTCCCAAGCCCTCGCCCACTGCTCCAGCAGCCAGCCCAACACACAGAGGGGACTG 195
QY 2990 ccactctcccttgcagctgctgagccgcagagagtgacggttctctcacagaggaggg 3049
Db 194 CCACCTCCCTTGGCAGCTGCTGAGCCGCAGAGAGTACAGGTTCTCTACACAGGACAGGG 135
QY 3050 gttcctttgggcatcatcagcagtagaatacaatttgggtgattggatctgtgt 3109
Db 134 GTTCTTCTGGGCATTACATCGCATAGAAATCAATAATTTGTGGTGATTGGATCTGTGT 75
QY 3110 tttaatgatttcagtgatttattgatttaattgtgcaagcttttctctaataaagc 3169
Db 74 TTTAATGAGTTTCAGTGATTTGATTGATTATTAATTGCAAGCTTTTCTCTAATAAAGC 15
QY 3170 tggagaatcaca 3181
Db 14 TGGAGAAATCACA 3

RESULT 12
AI439544
LOCUS          493 bp      mRNA      EST      28-MAR-1999
DEFINITION    tc90b05.x1 NCI_CGAP_CLL1 Homo sapiens cDNA clone IMAGE:2073393 3'
              similar to SW:CIK9_MOUSE P97414 VOLTAGE-GATED POTASSIUM CHANNEL
              PROTEIN KV1.9. [1] ; mRNA sequence.
ACCESSION     AI439544
VERSION       AI439544.1 GI:4305065
KEYWORDS      EST.
SOURCE        human.
ORGANISM      Homo sapiens
              Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
              Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE     1 (bases 1 to 493)
AUTHORS      NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE        National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
              Tumor Gene Index
```

JOURNAL
COMMENT

Unpublished (1997)

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-re@mail.nih.gov

Tissue Procurement: Ash Alizadeh, John Byrd, M.D., Mike Grever,

M.D., Louis M. Staudt, M.D., Ph.D.

cDNA Library Preparation: M. Bento Soares, Ph.D.

cDNA Library Arrayed by: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CGAP clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.llnl.gov/bbrp/image/image.html

Insert Length: 1584 Std Error: 0.00

Seq primer: -400P from Gibco

High quality sequence stop: 393.

FEATURES

source

Location/Qualifiers

1. 493

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:2073393"

/clone_lib="NCI_CGAP_CLL1"

/tissue_type="B-cell, chronic lymphocytic leukemia"

/lab_host="DH10B"

/note="Vector: pT7T3D-Pac (Pharmacia) with a modified

polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA

was primed with a Not I - oligo(dT) primer [5'

TGTTCACCAATCTGAAGTGGAGCGCGCATGCTTTTTTTTTTTTTTTTTTTT

T 3']; double-stranded cDNA was ligated to Eco RI

adaptors (Pharmacia), digested with Not I and cloned into

the Not I and Eco RI sites of the modified pT7T3 vector.

Library is normalized, and was constructed by Bento

Soares and M.Fatima Bonaldo."

64 a 168 c 151 g 110 t

BASE COUNT

ORIGIN

Query Match

Best Local Similarity 100.0%; Pred. No. 1.2e-209;

Matches 440; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 405 gcggcgccgcggtagacctagaccgcgcgtctccatctacagcagcgccgcgtgtt 464

Db 1 GCGGCGCGCGGTGAGCGCTAGACCGCGCGTCTCCATCTACAGCACGCGCGCGGTGT 60

QY 465 ggcgcgcacccacgtccagggcgcgctctacaacttctcgagcgtccacacggctgaa 524

Db 61 GCGGCGCACCCAGTCCAGGGCGCGCTCTACAACTTCTCGAGCGTCCACCGGCTGGAA 120

QY 525 atgctctgtttaccacttcgcgcttctctcatcgtctgtgtgtcctcatcttcagcgt 584

Db 121 ATGCTTCGTTTACCACCTTCGCGGTCTTCCTCATCGTCTCTGCTCATCTTCAGCGT 180

QY 585 gctgtccaccatcgagcaatgacgcgcctgcccacgggactctcttctgagatgagat 644

Db 181 GCTGTCCACATCGAGCAGATATGCGCGCCCTGCGCCAGCGGAGTCTCTTCTGGATGAGAT 240

QY 645 cgtgctgggtgttcttcgggacggagtagcgtggccgcctctggtcgcgcggcgctgcg 704

Db 241 CGTGTGTGGTGTCTTCGCGGACGGAGTACGTGTGCGGCTCTGTCGCGGCTGCCG 300

QY 705 cagcaagtacgtgggctctggggcggtgcgtcttggcccggaagcccatcttccatcat 764

Db 301 CAGCAAGTACGTGGGCGCTCTGGGGCGGCGTGCCTTTGCCCGGGAAGCCCATTTCCATCAT 360

QY 765 cgaactcatcgtggctgctgagcctcatcagtggtcctctcgtggtggtcgaagggcaggt 824

Db 361 CGACCTCATCGTGGTGGCGCTCCATGCTGTGCTCTCGTGGGCTCCAAAGGGGAGGT 420

QY 825 gttgccacgtcgccatca 844

Db 421 GTTTGCCAGTCGGCCATCA 440

RESULT 13

AW005483/c
LOCUS 484 bp mRNA 08-MAR-2000
DEFINITION w94f02.x1 NCI_CGAP.Co3 Homo sapiens cDNA clone IMAGE:2505627 3', similar to contains TAR1.tl TAR1 repetitive element ; , mRNA sequence.
ACCESSION AW005483
VERSION AW005483.1 GI:5854261
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 484)
REFERENCE NCI_CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
AUTHORS National Cancer Institute, Cancer Genome Anatomy Project (CGAP), TITLE Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Elias Campo, M.D., Michael R. Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D.
cDNA Library Arraying: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LENL at: www-bio.lnl.gov/bbrp/image/image.html
Insert length: 685 Std Error: 0.00
Seq primer: -40UP from Gibco
High quality sequence stop: 404.
FEATURES
source Location/Qualifiers
1..484
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2505627"
/clone_lib="NCI_CGAP_Co3"
/sex="pooled"
/tissue_type="colon"
/lab_host="DH10B"
/notes="Vector: pT73D-Pac (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA was prepared from 12 pooled bulk tumor samples and primed with a Not I - oligo(dT) primer. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT73 vector. Library went through one round of normalization."
BASE COUNT 108 a 125 c 141 g 110 t
ORIGIN
Query Match 12.8%; Score 408; DB 110; Length 484;
Best Local Similarity 100.0%; Pred. No. 1.4e-193;
Matches 408; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 2774 tgagcccccagctccagcaggagagcagctcacatttccccagggcacgtggtga 2833
Db 415 TGAGCCCCCAGCTTCCAGCAGAGGGACAGTCTCACCATTTCCTCCAGGGCAGGTGTTGA 356
Qy 2834 gtgagggggaacgcccacttccctgggttagactgccagcttcttcagctgagagagc 2893
Db 355 GTGGGGGGAACGCCACTTCCCTGGGTAGACTGCCAGCTCTCCTAGCTGGAGGAGC 296
Qy 2894 cctgcctctccgcccctgagcccactgtgcgtggggctccgcctccaaacccctcgccca 2953
Db 295 CCTGCCCTTCGCGCCCTTAGCCCACTGTGCGTGGGGGTCCCGCCTCCAAACCCCTCGCCCA 236
Qy 2954 gtccacgacgacgccaacacacagaaggagactgcacactccctctgcagctgctga 3013
Db 235 GTCCACGACGACGCCAAACACACAGAAGGGGACTGCCAGCTCCCTTGCAGCTGCTGA 176
Qy 3014 gccgcagagaagtgcaggttccctcacaggagacaggggttctcttggtggcattacatcgca 3073

Dd 175 GCCGACAGAGAAGTGACGGTTCCTACACAGGACAGGGGTTCCTCTCGGCATTAACATCGCA 116

Qy 3074 tagaaatcaataatttggtgattggatctgctgttttaagtgaagttccacagtgtgatt 3133
|||||
Dd 115 TAGAAATCAATAATTGTGGTGATTGGATCTGTGTTTTAATGAGTTTCACAGTGTGATT 56
|||||

Qy 3134 ttgattattaattgcgaagcttttctctaataaacgtggagaatcaca 3181
|||||
Dd .55 TTGATTATTAA'TGTGAAGCTTTTCCCTAATAA'ACG'TGGAGAAT'CACA 8
|||||

RESULT 14

BFI09081 451 bp mRNA EST 20-OCT-2000

LOCUS 715006.x1 Soares.NSF_F8_9W_OT_PA_P_S1 Homo sapiens cDNA clone

DEFINITION IMAGE:3524698 3', similar to SW-CIOL HUMAN P51787 VOLTAGE-GATED POTASSIUM CHANNEL PROTEIN KQT-LIKE 1 ; , mRNA sequence.

ACCESSION BFI09081

VERSION BFI09081.1 GI:10938851

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.

1 (bases 1 to 451)

NCI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index

JOURNAL Unpublished (1997)

CONTACT: Robert Strausberg, Ph.D.

EMAIL: rcapbs-r@mail.nih.gov

This clone is available royalty-free through LLNL ; contact the IMAGE Consortium (info@image.llnl.gov) for further information.

Seq primer: -40UP from Gibco

High quality sequence stop: 354.

Location/Qualifiers

1..451

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:3524698"

/clone_lfb="Soares.NSF_F8_9W_OT_PA_P_S1"

/lab_host="DH10B"

/notes="Organ: pooled; Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Site_1: Not 1; Site_2: Eco RI;

Equal amounts of plasmid DNA from five normalized libraries were mixed, and ss circles were made in vitro.

Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from pools of 5,000 clones made from the same 5 libraries. The pools consisted of the following libraries and clonesIDs: Soares NBHSF pool 1: 309384-310919, 323208-325895 Soares Nb2HP pool 1: 145032-147335, 147720-148103, 148872-149255, 15002 - 150407, 151176-152327 Soares Nb2HF8-9W pool 1: 75280-760583, 772104-774407 Soares NbHPA pool 1: 304776-306311, 320136-322823, 326280-326663 Soares NBHOT pool 1: 723720-726407, 739080-740999 Subtraction by Bento Soares and M. Fatima Bonaldo."

62 a 153 c 128 g 108 t

BASE COUNT

ORIGIN :

Query Match 12.4% Score 393; DB 144; Length 451;
Best Local Similarity 100.0%; Pred. NO. 5e-186;
Matches 393; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 423 agaccgcgcgtctccatctcacgacgcgcgcgcccggtgttgscgcgcaccccagtcaca 482
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Dd 1 AGACCCGCCGCTCTCCATCTACACAGCGCGCGCCGGTGTGGCGCGCACCCACGTCGA 60
|||||

Qy 483 ggccgcgcgtctacacctctctccagcgcgtcccccacgcggtcggaatgcttcgtttaccactt 542
|||||

Dd .61 GGCGCGGCTCTACAACTTCCTCGACGCTCCACCGGCTGGAATGCTTCGTTTACCACATT 120
|||||

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: November 2, 2001, 12:12:04 ; Search time 294.94 Seconds
(without alignments)
6772.072 Million cell updates/sec

Title: US-09-135-010A-1
Perfect score: 3181
Sequence: 1 ctgcccctccggcccccgc.....aataaacgtggagaatcaca 3181

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 730101 seqs, 313950809 residues

Word size : 12

Total number of hits satisfying chosen parameters: 106307

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : N_Geneseq_0601:*

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2: /SIDSI/gcgdata/geneseq/geneseq/NA1981.DAT:*
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22: /SIDSI/gcgdata/geneseq/geneseq/NA2001.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	3181	100.0	3181	21	Human KVLQT1 prote
2	3181	100.0	3181	21	Human long QT synd
3	3181	100.0	3181	22	Human KVLQT1 codin
4	3061	96.2	3182	22	Mutant human KVLQT
5	2702	84.9	2734	22	Mutant human KVLQT
6	2702	84.9	2821	18	DNA encoding human
7	2702	84.9	2821	18	Human KVLQT1 full-
8	265	8.3	494	21	Human pancreatic c
9	159	5.0	432	21	Human colon cancer
10	96	3.0	2821	18	DNA encoding human
11	96	3.0	2821	18	Human KVLQT1 full-

12	45	1.4	45	20	AAZ11946	Human potassium ch
13	41	1.3	83	16	AAT26420	Human gene signatu
14	34	1.1	2734	22	AAC89984	Mutant human KVLQT
15	22	0.7	22	18	AAT91065	Human KVLQT1 S2-S3
16	22	0.7	22	18	AAT90717	Human KVLQT1 S2-S3
17	22	0.7	22	21	AAZ90741	Human KVLQT1 mutat
18	22	0.7	22	21	AAZ98971	Mutant human long
19	21	0.7	2335	21	AAA47618	KCNQ4 Potassium ch
20	21	0.7	910715	20	AAZ20248	Borrelia burgdorfe
21	20	0.6	20	18	AAT91069	Human KVLQT1 S4 re
22	20	0.6	20	18	AAT90721	Human KVLQT1 S4 re
23	20	0.6	20	21	AAZ90745	Human KVLQT1 mutat
24	20	0.6	20	21	AAZ98975	Mutant human long
25	20	0.6	936	20	AAZ56375	Human DNA-dependen
26	20	0.6	2065	19	AAV29062	BRCA1 modulator pr
27	20	0.6	2065	20	AAZ86754	CDNA 091-21A31 enc
28	19	0.6	19	21	AAZ90707	Forward primer for
29	19	0.6	19	21	AAZ90708	Reverse primer for
30	19	0.6	19	21	AAZ98937	Human long QT synd
31	19	0.6	19	21	AAZ98938	Human long QT synd
32	19	0.6	19	22	AAC89947	Human KVLQT1 exon
33	19	0.6	19	22	AAC89948	Human KVLQT1 exon
34	19	0.6	263	21	AAC24694	Human secreted pro
35	19	0.6	770	20	AAV88894	EST clone H2162.
36	19	0.6	1182	21	AAA27105	Human h-TRAAK CDNA
37	19	0.6	1218	21	AAA27106	Human h-TRAAK CDNA
38	19	0.6	1323	21	AAC74299	Human secreted pro
39	19	0.6	1678	21	AAC99026	Human pancreatic c
40	19	0.6	1765	22	AAC93864	Human CDNA encodin
41	19	0.6	2786	16	AAZ06023	CDNA encoding aven
42	19	0.6	3269	16	AAZ06024	CDNA encoding aven
43	19	0.6	4132	20	AAZ60265	Nucleic acid seque
44	19	0.6	6910	21	AAA38335	Human aldosterone
45	19	0.6	7011	19	AAV20464	Human L-myc oncoge

ALIGNMENTS

RESULT 1

AAZ90669

ID AAZ90669 standard; cDNA; 3181 BP.

AC AAZ90669;

DT 19-JUN-2000 (first entry)

XX Human KVLQT1 protein encoding cDNA.

DE KVLQT1; KCNE1; long QT syndrome; LQT syndrome; mink protein;

KW antiarrhythmic; gene therapy; human; ss.

XX Homo sapiens.

OS

XX Key

FH Location/Qualifiers

FT 163..2193

FT /*tag= a

FT /product= "KVLQT1"

XX

PN WO200006600-A1.

XX

PD 10-FEB-2000.

XX

PF 06-OCT-1998; 98WO-US17838.

XX

PR 29-JUL-1998; 98US-0094477.

PR 17-AUG-1998; 98US-0135020.

XX

PA (UTAH) UNIV UTAH RES FOUND.

XX

PI Ksating MT Sanguinetti MC, Splawski I;

DR WPI; 2000-195262/17.

DR P-PSDB; AAY57368.

XX Mutant forms of genes encoding minK protein and KVLOT1 protein involved
PT in cardiac potassium channel formation useful for screening drugs, for
PT preventing and treating cardiac arrhythmia -
XX
XX
PS Claim 28; Fig 5A-B; 167pp; English.

XX The invention relates to KVLOT1 and KCNE1 genes, associated with long
CC QT (LQT) syndrome. It provides a minK protein comprising a mutation which
CC substitutes the wild type amino acids with Leu, Asp, Leu, His, Trp and
CC Ala or Thr at residues 74, 76, 28, 32, 98 and 127 respectively. Screening
CC KVLOT1 and KCNE1 is useful for identifying mutations for diagnosing and
CC treating LQT. The ability to predict LQT enables physicians to prevent
CC the diseases with medical therapy such as beta blocking agents and ops
CC for better treatments. The present sequence represents the cDNA encoding
CC the human KVLOT1 protein.

XX Sequence 3181 BP; 581 A; 1073 C; 968 G; 559 T; 0 other;

Query Match 100.0%; Score 3181; DB 21; Length 3181;

Best Local Similarity 100.0%; Pred. No. 0;

Matches 3181; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ctgccccctccggccccggccgagcgccccgggtggcgccgagcgcccccgcgcg 60

Db 1 ctgccccctccggccccggccgagcgccccgggtggcgccgagcgcccccgcgcg 60

QY 61 gggctggcagcagtggtgctccgcagctcgccccggcgctcgcttcgctgcagctcccg 120

Db 61 gggctggcagcagtggtgctccgcagctcgccccggcgctcgcttcgctgcagctcccg 120

QY 121 gtgcgcgcgtcggtggccccggccccggcgccctctgttatggcgcggtcctcc 180

Db 121 gtgcgcgcgtcggtggccccggccccggcgccctctgttatggcgcggtcctcc 180

QY 181 ccgccccggcgaggaagcgtgggttgggggcgccgctgcagcgccccggcggggc 240

Db 181 ccgccccggcgaggaagcgtgggttgggggcgccgctgcagcgccccggcggggc 240

QY 241 agcgcgccgtgacaaagtgccctctcgctgagctgagctggcgagggcgccggcg 300

Db 241 agcgcgccgtgacaaagtgccctctcgctgagctgagctggcgagggcgccggcg 300

QY 301 ggcgcgcgtctacgccccatcgccccggcgccccaggtcccgccccctgcgtcc 360

Db 301 ggcgcgcgtctacgccccatcgccccggcgccccaggtcccgccccctgcgtcc 360

QY 361 ccggcgcgcccgccccagttgctctccgaccttgcccgcgccggcgccggtgagc 420

Db 361 ccggcgcgcccgccccagttgctctccgaccttgcccgcgccggcgccggtgagc 420

QY 421 ctgagccgcgctctccatctacagcagcgccccggcggtgttgcgcgccaccagtc 480

Db 421 ctgagccgcgctctccatctacagcagcgccccggcggtgttgcgcgccaccagtc 480

QY 481 caggccgcgtctacaacttctcagcgctccaccggcggtgaaatgcttcgtttaccac 540

Db 481 caggccgcgtctacaacttctcagcgctccaccggcggtgaaatgcttcgtttaccac 540

QY 541 ttgcgcgttctctcagctggtgtctgctcctatcttccagcggtgttccaccatcgag 600

Db 541 ttgcgcgttctctcagctggtgtgtctcctatcttccagcggtgttccaccatcgag 600

QY 601 cagtatgcgcctggccacgggagctctctctggatggagatcgtggtgttc 660

Db 601 cagtatgcgcctggccacgggagctctctctggatggagatcgtggtgttc 660

QY 661 ttgggacgagtagtggttcgctctggttcgccccggtcgccgagcaagtacgtggcg 720

Db 661 ttgggacgagtagtggttcgctctggttcgccccggtcgccgagcaagtacgtggcg 720

QY 1801 caggggccacctcaacctcatgggtgcgcacatcaaggagctgcagaggaggtgaccagtc 1860

Db 1801 caggggccacctcaacctcatgggtgcgcacatcaaggagctgcagaggaggtgaccagtc 1860

QY 721 ctctggggcgcgctgctgttgcggagccatttccatcatcagacctcatcgtggtc 780

Db 721 ctctggggcgcgctgctgttgcggagccatttccatcatcagacctcatcgtggtc 780

QY 781 gtggcctccatggttggtcctctggtgggtcccaaggcgaggtgttgccacgtcgcc 840

Db 781 gtggcctccatggttggtcctctggtgggtcccaaggcgaggtgttgccacgtcgcc 840

QY 841 atcaggggatccctcttctcagatcctgaggtgctacacgtcgaccgcccaggagc 900

Db 841 atcaggggatccctcttctcagatcctgaggtgctacacgtcgaccgcccaggagc 900

QY 901 acctggaggtccctgggtcctggtttcctccaccgcccaggagctgataaccacctg 960

Db 901 acctggaggtccctgggtcctggtttcctccaccgcccaggagctgataaccacctg 960

QY 961 tacatcggttccctgggctcctatcttccctgacttctgtacctggtgagaagac 1020

Db 961 tacatcggttccctgggctcctatcttccctgacttctgtacctggtgagaagac 1020

QY 1021 gcggtgaacgagtcagcgcggtggttcggcagctacgcagatgcgtgtggtgggg 1080

Db 1021 gcggtgaacgagtcagcgcggtggttcggcagctacgcagatgcgtgtggtgggg 1080

QY 1081 gtggtcagtcacccaccatcggtatggggacaaagtgccccacagctggtgcggaa 1140

Db 1081 gtggtcagtcacccaccatcggtatggggacaaagtgccccacagctggtgcggaa 1140

QY 1141 acctgcctcctctctctctgttcttggcctctctctcttcttgcgtcccggtgatt 1200

Db 1141 acctgcctcctctctctctgttcttggcctctctctcttcttgcgtcccggtgatt 1200

QY 1201 cttggtcgggtttgcccagaaagtgccagaaagcagagcagaagcaacttcaacgg 1260

Db 1201 cttggtcgggtttgcccagaaagtgccagaaagcagagcagaagcaacttcaacgg 1260

QY 1261 cagatcccgcgcgagcctcactcattcagaccgctgaggtgctatgctgcgagaac 1320

Db 1261 cagatcccgcgcgagcctcactcattcagaccgctgaggtgctatgctgcgagaac 1320

QY 1321 ccgactcctccactggaagatctacatccgaaagccccccggagccacactctgtg 1380

Db 1321 ccgactcctccactggaagatctacatccgaaagccccccggagccacactctgtg 1380

QY 1381 tcacccgccccaaacccaaagctctgtgtgtaaaagaaaaaaagtccaagtgcac 1440

Db 1381 tcacccgccccaaacccaaagctctgtgtgtaaaagaaaaaaagtccaagtgcac 1440

QY 1441 aaagacaatgggtgactcctggagagaagatgctcacagtcctcccatatcacgtgcac 1500

Db 1441 aaagacaatgggtgactcctggagagaagatgctcacagtcctcccatatcacgtgcac 1500

QY 1501 cccccaagagcgcgctgagccacttctctgcagcgctatgacagttctgtaagg 1560

Db 1501 cccccaagagcgcgctgagccacttctctgcagcgctatgacagttctgtaagg 1560

QY 1561 aagagcccaactgctgggaagtgcacatcccatcttcagagaaacacagcttcgcc 1620

Db 1561 aagagcccaactgctgggaagtgcacatcccatcttcagagaaacacagcttcgcc 1620

QY 1621 gaggacctggacctggaggggagactctgacacccatcacccatctcacagctg 1680

Db 1621 gaggacctggacctggaggggagactctgacacccatcacccatctcacagctg 1680

QY 1681 cgggaacacatcgggccacatttaagtgatctgcagcgtacgtacttctggtgccaag 1740

Db 1681 cgggaacacatcgggccacatttaagtgatctgcagcgtacgtacttctggtgccaag 1740

QY 1741 aaaaaatccagaaagcggaagccttgcagcgtacgtacttctggtgccaag 1800

Db 1741 aaaaaatccagaaagcggaagccttgcagcgtacgtacttctggtgccaag 1800

QY 1801 caggggccacctcaacctcatgggtgcgcacatcaaggagctgcagaggaggtgaccagtc 1860

Db 1801 caggggccacctcaacctcatgggtgcgcacatcaaggagctgcagaggaggtgaccagtc 1860

```

Db 1801 cagggccactcaacctcaatagtcgcgcaataaagagctgcagagagagctggaccagctcc 1860
Qy 1861 attgggaagccctcaactgtttcatctccgtctcagaaaaagacaaagatcgcggcgagcaac 1920
Db 1861 attgggaagccctcaactgtttcatctccgtctcagaaaaagacaaagatcgcggcgagcaac 1920
Qy 1921 acgatacggcgcgcctgaaacagagtagaagaacaaggtgacgcagctggaccagaggtg 1980
Db 1921 acgatacggcgcgcctgaaacagagtagaagaacaaggtgacgcagctggaccagaggtg 1980
Qy 1981 gcaactcalcaccagacatgcttcacagctgctctccttgccacggtgacgcagacccccgcgc 2040
Db 1981 gcaactcalcaccagacatgcttcacagctgctctccttgccacggtgacgcagacccccgcgc 2040
Qy 2041 agcggcgcccccagagagggcgccacacatcacccagccctgcggcagtgccggc 2100
Db 2041 agcggcgcccccagagagggcgccacacatcacccagccctgcggcagtgccggc 2100
Qy 2101 tccgtcacccctgagctcttctgcccacagcaacacccctgcacacacacagcagctgacc 2160
Db 2101 tccgtcacccctgagctcttctgcccacagcaacacccctgcacacacacagcagctgacc 2160
Qy 2161 gtcgccagagggggcccgatcgaggggtcctcgaggaggaggtgggctgggggatggggcc 2220
Db 2161 gtcgccagagggggcccgatcgaggggtcctcgaggaggaggtgggctgggggatggggcc 2220
Qy 2221 tgagtgagagggggagccaaagtgggcccccacccctgcccctctctgaagagggccacccctcc 2280
Db 2221 tgagtgagagggggagccaaagtgggcccccacccctgcccctctctgaagagggccacccctcc 2280
Qy 2281 taaaagcccagagagagagagagagagagagagagagagagagagagagagagagagagagag 2340
Db 2281 taaaagcccagagagagagagagagagagagagagagagagagagagagagagagagagagag 2340
Qy 2341 gtcggcacagctgcacttgagggtgcagcagcagcagcagcagcagcagcagcagcagcagc 2400
Db 2341 gtcggcacagctgcacttgagggtgcagcagcagcagcagcagcagcagcagcagcagcagc 2400
Qy 2401 gcccctctcaggtctgagtggttaccccaagcgcctggccccacacatggtgatgttga 2460
Db 2401 gcccctctcaggtctgagtggttaccccaagcgcctggccccacacatggtgatgttga 2460
Qy 2461 catcactggcatggtgtgtggaccacagtgacagggcagggcagggcagggcagggcagggc 2520
Db 2461 catcactggcatggtgtgtggaccacagtgacagggcagggcagggcagggcagggcagggc 2520
Qy 2521 caggaaatagcacagggctgagtgagggcccccacccctgcttgcccaggggggttctctgagg 2580
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Qy 2581 ggagacagagcaacccctggaccacagcctcaaatccaggacccctgcaggacacagggcag 2640
Db 2581 ggagacagagcaacccctggaccacagcctcaaatccaggacccctgcaggacacagggcag 2640
Qy 2641 ggaggaccagccacagctgactacagggccacggcgaataaaagcccgagagccattt 2700
Db 2641 ggaggaccagccacagctgactacagggccacggcgaataaaagcccgagagccattt 2700
Qy 2701 ggaggccctgggctggtcctcactctcaggaatgctgacccatgggcagggagactg 2760
Db 2701 ggaggccctgggctggtcctcactctcaggaatgctgacccatgggcagggagactg 2760
Qy 2761 tggagactgctctgagccccagcttccagcagagaggagagagagagagagagagagagagag 2820
Db 2761 tggagactgctctgagccccagcttccagcagagaggagagagagagagagagagagagagag 2820
Qy 2821 ggcagctggttgagtgagggggaagccacacttccctgggttagagctgcccagcttctcta 2880
Db 2821 ggcagctggttgagtgagggggaagccacacttccctgggttagagctgcccagcttctcta 2880
Qy 2881 gctggagagagccctgctctccgcctcagccccactgctgctgggggctcccgccctcc 2940
Db 2881 gctggagagagccctgctctccgcctcagccccactgctgctgggggctcccgccctcc 2940
```

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Db 2881 gctggagagagagccctgctctccgcctcagccccactgctgctgggggctcccgccctcc 2940
Qy 2941 aacccctcgcagctccagcagccagccacacacacacacacacacacacacacacacacacac 3000
Db 2941 aacccctcgcagctccagcagccagccacacacacacacacacacacacacacacacacacac 3000
Qy 3001 tgccagctgctgagcgcgcagagaaagtgcagcgttccctacacagagacaggggttctcttgg 3060
Db 3001 tgccagctgctgagcgcgcagagaaagtgcagcgttccctacacagagacaggggttctcttgg 3060
Qy 3061 gcaattacatcgatagaataatattgtgtgtgattggtgattggtgattggtgattggtgattg 3120
Db 3061 gcaattacatcgatagaataatattgtgtgtgattggtgattggtgattggtgattggtgattg 3120
Qy 3121 tcacagctgatttattattattattattattattattattattattattattattattattatt 3180
Db 3121 tcacagctgatttattattattattattattattattattattattattattattattattatt 3180
Qy 3181 a 3181
Db 3181 a 3181

RESULT 2
AAZ98901
ID AAZ98901 standard; cDNA; 3181 BP.
XX
AC AAZ98901;
XX
DT 06-JUN-2000 (first entry)
XX
DE Human long QT syndrome-associated KVLQT1 cDNA.
KW KVLQT1; mutation; human; cardiac I(ks) potassium channel; KCNE1; ss;
KW cardiac arrhythmia; electrocardiogram; Long QT syndrome; gene therapy;
KW chromosome 11p15.5.
XX
OS Homo sapiens.
XX
PN WO200006199-A1.
XX
PD 10-FEB-2000.
XX
PF 12-MAY-1999; 99WO-US10260.
XX
PR 29-JUL-1998; 98US-0094477.
PR 17-AUG-1998; 98US-0135010.
XX
PA (UTAH ) UNIV UTAH RES FOUND.
PA (GENZ ) GENZYME CORP.
XX
PI Seating, MT Sanguinetti MC, Curran ME, Landes GM, Connors TD;
PI Burn TC, Splawski I;
XX
DR WPI; 2000-195199/17.
DR P-PSDB; AAY80562.
XX
PT New isolated mutant KVLQT1 nucleic acids, useful for developing
PT products for the diagnosis, prevention and treatment of long QT
PT syndrome -
XX
PS Claim 1; Fig 5A-B; 178pp; English.
XX
CC The invention relates to KVLQT1 nucleic acids which have a mutation
CC compared to wild-type KVLQT1. This sequence corresponds to the human
CC wild type KVLQT1 cDNA sequence. The sequence encodes a protein of 676
CC amino acid which forms a cardiac I(ks) potassium channel with the KCNE1
CC protein (AAY80563). The KVLQT1 gene contains 15 introns and encodes a
CC protein containing 6 putative transmembrane segments and a pore forming
CC region. The gene has been mapped to the chromosomal location 11p15.5.
CC Mutations in the KVLQT1 or KCNE1 genes result in cardiac arrhythmias
CC observed as a prolonged QT curve in electrocardiograms (Long QT
CC syndrome). The genes and proteins can be used for the diagnosis of
```


Ov 121 ataccac

Qy	1	ctgcccctcggccccgcgccccgagcgcgcggctggccggcagcagcccccgcgcg	60
Db	1	ctgcccctcggccccgcgccccgagcgcgcggctggccggcagcagcccccgcgcg	60
Qy	61	ggctggcagcagtgctcccgcactcgccccggcgtcgccccggcgtcgcttcgctcagctcccg	120
Db	61	ggctggcagcagtgctcccgcactcgccccggcgtcgccccggcgtcgcttcgctcagctcccg	120
Qy	121	gtgcgcgcgtcgggcggccccccggcaggccctccctcgttatggcgcgggcctcctcc	180

CC The present sequence is the coding sequence for wild-type human KVLOQT1.
CC KVLOQT1 is a cardiac potassium channel and mutations in the KVLOQT1 gene
CC cause Jervell and Lange-Nielsen Syndrome (JLN). KVLOQT1 maps to
CC chromosome 1p15.5. The present invention relates to a mutant KVLOQT1
CC coding sequence (see AAC89914). The mutant KVLOQT1 coding sequence is
CC useful in the diagnosis of long QT syndrome and in screening humans for
CC the presence of KVLOQT1 gene variants which cause JLN syndrome.

XX
SQ Sequence 3181 BP; 581 A; 1073 C; 968 G; 559 T; 0 other;


```
QY 2341 gctcggcacagccctgacacttgggggctcagcaaggccacactctctctgagccggtgtggg 2400
Db 2341 gctcggcacagccctgacacttgggggctcagcaaggccacactctctctgagccggtgtggg 2400
QY 2401 gccctcgtcaggtctgagttgtaccaccaagcgcctggccccacacatggtgatgtga 2460
Db 2401 gccctcgtcaggtctgagttgtaccaccaagcgcctggccccacacatggtgatgtga 2460
QY 2461 caccactggcaatggtgtgggacccagtgaggggacaggggctggccccatgtatggc 2520
Db 2461 caccactggcaatggtgtgggacccagtgaggggacaggggctggccccatgtatggc 2520
QY 2521 caggaaatagcacaggtgagtgagggcccaacctgctgtggccccagggggttcttgagg 2580
Db 2521 caggaaatagcacaggtgagtgagggcccaacctgctgtggccccagggggttcttgagg 2580
QY 2581 ggagacagagcaacccctggagcccgactcaaatccaggacctgcccagcacagggcag 2640
Db 2581 ggagacagagcaacccctggagcccgactcaaatccaggacctgcccagcacagggcag 2640
QY 2641 ggcaggaccgcccactgactacagggccacccggccaataaagccagagccattt 2700
Db 2641 ggcaggaccgcccactgactacagggccacccggccaataaagccagagccattt 2700
QY 2701 ggaaggcctgggctgctccctcactctcaggaataatgctgacccatgggcaggaactg 2760
Db 2701 ggaaggcctgggctgctccctcactctcaggaataatgctgacccatgggcaggaactg 2760
QY 2761 tgagactgctcctgagccccagcttcacagagggagacagtcacatttcccag 2820
Db 2761 tgagactgctcctgagccccagcttcacagagggagacagtcacatttcccag 2820
QY 2821 ggcacgtggttggagggggaaagccacacttccctgggttagactgacagctcttcta 2880
Db 2821 ggcacgtggttggagggggaaagccacacttccctgggttagactgacagctcttcta 2880
QY 2881 gctgagagagccctgctcctcgcctcagccactgagccactgctggtgggctcccgctcc 2940
Db 2881 gctgagagagccctgctcctcgcctcagccactgagccactgctggtgggctcccgctcc 2940
QY 2941 aacccctcgccagctccagcagcagcagcacaacacagaggggactgcccactccct 3000
Db 2941 aacccctcgccagctccagcagcagcagcacaacacagaggggactgcccactccct 3000
QY 3001 tgcagctgctgagccgacagagagtgacggttctctacacagagaggggttctctgg 3060
Db 3001 tgcagctgctgagccgacagagagtgacggttctctacacagagaggggttctctgg 3060
QY 3061 gcattacacgcataagaatacaataattgtggtgatttgatctgtgttttaagatt 3120
Db 3061 gcattacacgcataagaatacaataattgtggtgatttgatctgtgttttaagatt 3120
QY 3121 tcacagtgatttattattattgtgcaagcttttccataaataaagtgagagataac 3180
Db 3121 tcacagtgatttattattattgtgcaagcttttccataaataaagtgagagataac 3180
QY 3181 a 3181
Db 3181 a 3181
```

RESULT 4

```
AAC89914
ID AAC89914 standard; cDNA; 3182 BP.
XX AC
XX AC AAC89914;
XX AC
XX 08-MAR-2001 (first entry)
XX Mutant human KVLQT1 coding sequence #2.
XX Human; KVLQT1; antiarrhythmic; cardiant; gene therapy;
KW cardiac potassium channel; Jervell and Lange-Nielsen Syndrome; JLN;
```

```
KW chromosome 11p15.5; long QT syndrome; ss.
OS Homo sapiens.
XX US6150104-A.
PD 21-NOV-2000.
XX 17-AUG-1998; 98US-0135021.
XX 29-JUL-1998; 98US-0094477.
PR 13-JUN-1997; 97US-0874655.
XX (UTAH ) UNIV UTAH RES FOUND.
PA (Keating MT) Splawski I;
PI WPI: 2001-060013/07.
DR P-PSDB; AAB49495.
XX DNA encoding for a mutant KVLQT1 which causes Jervell and Lange-Nielsen
syndrome (JLN) when homozygous, useful for diagnosing long QT syndrome,
or diagnosing or prognosing JLN -
XX Example 4; Columns 63-68; 58pp; English.
XX KVLQT1 is a cardiac potassium channel and mutations in the KVLQT1 gene
cause Jervell and Lange-Nielsen Syndrome (JLN). KVLQT1 maps to
CC chromosome 11p15.5. The present sequence is a mutant KVLQT1 coding
CC sequence. The mutant KVLQT1 coding sequence is useful in the diagnosis of
CC long QT syndrome and in screening humans for the presence of KVLQT1 gene
CC variants which cause JLN syndrome.
XX Sequence 3182 BP; 581 A; 1073 C; 969 G; 559 T; 0 other;
```

```
Query Match 96.2%; Score 3061; DB 22; Length 3182;
Best Local Similarity 100.0%; Pred. No. 0;
Matches 3181; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY 1 ctgccccctcggcccgcccgagcgccggctggcgccgagcgcccgcccgcccgccg 60
Db 1 ctgccccctcggcccgcccgagcgccggctggcgccgagcgcccgcccgcccgccg 60
QY 61 gggctggcagcagtgctgctccgcacactgcccggcgctcgccttcgctgagctcccg 120
Db 61 gggctggcagcagtgctgctccgcacactgcccggcgctcgccttcgctgagctcccg 120
QY 121 gtgcgcgcgctcggcgccggcccccggcagggccctcctctgttatggcgcgccctcc 180
Db 121 gtgcgcgcgctcggcgccggcccccggcagggccctcctctgttatggcgcgccctcc 180
QY 181 ccgcccagggcgagagagagcgctgggttggggcgccctgcccagggcccgccgggggc 240
Db 181 ccgcccagggcgagagagagcgctgggttggggcgccctgcccagggcgcccgccgggc 240
QY 241 agcgcgcgctggccaagaagtgcctctcctcgtcagctgagctgagggcgcccgccg 300
Db 241 agcgcgcgctggccaagaagtgcctctcctcgtcagctgagggcgcccgcccgccg 300
QY 301 ggcgcgcgcgtctacgcgcacatgcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgc 360
Db 301 ggcgcgcgcgtctacgcgcacatgcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgc 360
QY 361 ccggcgcccgcccgcccgcccgagttgcctccgaaccttggccccgcgcccgccggtagc 420
Db 361 ccggcgcccgcccgcccgcccgagttgcctccgaaccttggccccgcgcccgccggtagc 420
QY 421 ctgagcccgcgcttccatctacacagcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgc 480
Db 421 ctgagcccgcgcttccatctacacagcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgc 480
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DB 1113 gaagagcccaacactctgtgaagtgaatgcgccatttcatgagaacaaacagctcgc 1172
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QY 3180 ca 3181
DB 2733 ca 2734

RESULT 6

AAT94004
ID AAT94004 standard; DNA; 2821 BP.

XX AC AAT94004;

XX DT 28-FEB-1998 (first entry)

XX DE DNA encoding human KVLQT1.

XX KW KVLQT1; long QT syndrome; arrhythmia; minK; potassium channel;
diagnosis; therapy; human; ds.

XX OS Homo sapiens.

XX FT Key Location/Qualifiers
CDS 88..1833

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FT XX      /*tag= a
PN XX      WO9723632-A1.
PD XX      03-JUL-1997.
PF XX      20-DEC-1996; 96WO-US19917.
PS XX      29-OCT-1996; 96US-0739383.
PR XX      22-DEC-1995; 95US-0019014.
PA XX      (GEN2 ) GENZYME GENETICS.
PA XX      (UTAH ) UNIV UTAH RES FOUND.
XX XX      Connors TD, Curran ME, Keating MF, Landes GM;
XX XX      WPI; 1997-402191/37.
XX XX      P-PSDB; AAW33355.
XX XX      New isolated human potassium channel gene, KVLQT1, - used to develop
XX XX      products for diagnosis, prevention and therapy of long QT syndrome
XX XX      Clalim 2; Page 76-79; 117pp; English.
XX XX      This cDNA sequence includes a full-length coding sequence for human
XX XX      KVLQT1 (see AAW33355), a novel cardiac potassium channel protein.
XX XX      The sequence was assembled from partial clones isolated from human
XX XX      pancreatic and cardiac cDNA libraries. KVLQT1 was mapped to
XX XX      chromosome 11p15.5 making it a candidate for the long QT syndrome
XX XX      (LQT) gene. LQT is an inherited cardiac arrhythmia. 16 Families
XX XX      with mutations in KVLQT1 have been identified and it was shown that
XX XX      in all 16 families there was complete linkage between LQT1 and
XX XX      KVLQT1. The KVLQT1 gene product coassembles with human mink to
XX XX      form a cardiac IKs potassium channel. IKs dysfunction is a cause
XX XX      of cardiac arrhythmia. Coexpression of KVLQT1 and mink in a host
XX XX      cell provides a means for screening for drugs useful in treating or
XX XX      preventing LQT. Analysis of the KVLQT1 gene will provide an early
XX XX      diagnosis of subjects with LQT. A claimed method of assessing the
XX XX      risk in a human for LQT syndrome comprises screening for a mutation
XX XX      in the KVLQT1 gene. Transgenic animals expressing human mink and
XX XX      KVLQT1 can be used to test therapeutic agents against LQT.
XX XX      Sequence 2821 BP; 562 A; 892 C; 843 G; 524 T; 0 other;
XX XX
XX XX      Query Match      84.9%; Score 2702; DB 18; Length 2821;
XX XX      Best Local Similarity 100.0%; Pred. NO. 0;
XX XX      Matches 2702; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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DB 240 gcagtagccgcctcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgcgc 299
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RESULT 7
AAT90730
ID AAT90730 standard; cDNA; 2821 BP.
XX
AC AAT90730;
XX
DT 12-FEB-1998 (first entry)
XX
DE Human KVLQTL full-length cDNA.
XX
KW KVLQTL; long QT syndrome; arrhythmia; minK; potassium channel;
diagnosis; therapy; human; ds.
XX
OS Homo sapiens.
XX
FH Key
FT CDS 88..1833
FT /*tag= a
XX
PN W09723598-A2.
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PD 03-JUL-1997.
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PF 20-DEC-1996; 96WO-US19756.
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PR 29-OCT-1996; 96US-0739383.
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PR 22-DEC-1995; 95US-0019014.
XX
XX (UTAH) UNIV UTAH RES FOUND.
XX Curran ME, Keating MT, Sanguinetti MC;
WPI; 1997-402190/37.
P-PSDB; AAW30038.
PT Human minK and Xenopus KVLQTL coding sequences - used for assays for
identifying drugs which can be used for preventing or treating long
QT syndrome
XX
PS Example 9; Page 76-79; 105pp; English.
XX
CC This cDNA sequence includes a full-length coding sequence for human
KVLQTL (see AAW30038), a novel cardiac potassium channel protein.
CC The sequence was assembled from partial clones isolated from human
pancreatic and cardiac cDNA libraries. KVLQTL was mapped to
CC chromosome 11p15.5 making it a candidate for the long QT syndrome
CC (LQT) gene. LQT is an inherited cardiac arrhythmia. One intragenic
CC deletion and 10 different missense mutations which cause LQT have
CC been identified in KVLQTL. The KVLQTL gene product coassembles
CC with human minK to form a cardiac IKs potassium channel.
CC Coexpression of these 2 proteins in a host cell provides a means
CC for screening for drugs useful in treating or preventing LQT. The
CC products can also be used for studying mechanisms underlying common
CC arrhythmias and for presymptomatic diagnosis of LQT. Transgenic
CC animals expressing human minK and KVLQTL can be used to test
CC therapeutic agents against LQT.
XX
SQ Sequence 2821 BP; 562 A; 892 C; 843 G; 524 T; 0 other;

Query Match		84.9%;	Score 2702;	DB 18;	Length 2821;
Best Local Similarity		100.0%;	Pred. No. 0;		
Matches 2702;		Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;
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DB	120	ccagggccgcgtctacaaactctcagagctccacacggtcgaaatgcttcgtttacca	179		
QY	540	cttcgcgctcttcctacgtctcgtggtctcgtctcgtctcgtctcgtctcgtctcgt	599		
DB	180	cttcgcgctcttcctacgtctcgtggtctcgtctcgtctcgtctcgtctcgtctcgt	239		
QY	600	gcagtatgcgcgcctggccacgggagctctctctctgagatgagatgctggtggtt	659		
DB	240	gcagtatgcgcgcctggccacgggagctctctctctgagatgagatgctggtggtt	299		
QY	660	cttcgggacagagtagctggtcgcgtctcgtcgcgtcgcgtcgcgtcgcgtcgcgt	719		
DB	300	cttcgggacagagtagctggtcgcgtctcgtcgcgtcgcgtcgcgtcgcgtcgcgt	359		
QY	720	cctctggggcgctgcgctcttgcgcggaagccatttccatcatgacctcatctggtt	779		
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QY	780	cgctggcctccatggtggtcctctcgttgggtcctcgaaggcgagtggttgcacgtcgc	839		
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QY	840	catcaggggacatccgtctcgtcagatcctcgtgagatgctacacgtcagcccgagg	899		
DB	480	catcaggggacatccgtctcgtcagatcctcgtgagatgctacacgtcagcccgagg	539		
QY	900	cacctgagagctcctggtccgttgcgtctcctcaccacgcagagagctgataaccacct	959		
DB	540	cacctgagagctcctggtccgttgcgtctcctcaccacgcagagagctgataaccacct	599		
QY	960	gtacatcggtctcctgggctcctctcctcgttacttctgtaacttctgtaacttctgta	1019		
DB	600	gtacatcggtctcctgggctcctctcctcgttacttctgtaacttctgtaacttctgta	659		
QY	1020	cgcggtgaacagatcagcgcggttggagttcggcagatcagcagatcgtctggtgggg	1079		
DB	660	cgcggtgaacagatcagcgcggttggagttcggcagatcagcagatcgtctggtgggg	719		
QY	1080	ggtggtcacagtcacacacatcgctatgggacaaagtgccacagcgtgggtcgggaa	1139		
DB	720	ggtggtcacagtcacacacatcgctatgggacaaagtgccacagcgtgggtcgggaa	779		
QY	1140	gacctgcgctcctgcttctctctctctcctcctcctcctcctcctcctcctcctcct	1199		
DB	780	gacctgcgctcctgcttctctctctctcctcctcctcctcctcctcctcctcctcct	839		
QY	1200	tcttgctcgggttggctcctgaagtgagcagcagcagcagcagcagcagcagcagcag	1259		
DB	840	tcttgctcgggttggctcctgaagtgagcagcagcagcagcagcagcagcagcagcag	899		
QY	1260	gcagatcccgccggcagcctcactcatctcagacgcagatgagagtgctatgctgcgagaa	1319		
DB	900	gcagatcccgccggcagcctcactcatctcagacgcagatgagagtgctatgctgcgagaa	959		
QY	1320	ccccgactcctccacctggagatctacatccggaagcccccgagccacacactctgct	1379		
DB	960	ccccgactcctccacctggagatctacatccggaagcccccgagccacacactctgct	1019		
QY	1380	gtcacccagcccccaaccccaagagtgctggtggttaaaagaaaaaaagttcaagctgga	1439		
DB	1020	gtcacccagcccccaaccccaagagtgctggtggttaaaagaaaaaaagttcaagctgga	1079		
QY	1440	caagagacaaatgggggtgactcctggagagagagatgctcacagtcctccatcatcagtcgca	1499		
DB	1080	caagagacaaatgggggtgactcctggagagagagatgctcacagtcctccatcatcagtcgca	1139		

QY	1500	cccccaagaagagcgcggtggaaccacttctctgacacggtctatgacagttctgtaag	1559		
DB	1140	cccccaagaagagcgcggtggaaccacttctctgacacggtctatgacagttctgtaag	1199		
QY	1560	gaagagcccaacactgctggaagtgagcatgcccatttcatgagaacaaacagcttcgc	1619		
DB	1200	gaagagcccaacactgctggaagtgagcatgcccatttcatgagaacaaacagcttcgc	1259		
QY	1620	cgagacactgacctggaggggagactctgctgacacccatcaccacatctcagct	1679		
DB	1260	cgagacactgacctggaggggagactctgctgacacccatcaccacatctcagct	1319		
QY	1680	gcgggaacacacatcgccacacatttaaggtctatcgagcgtatgcagttctggtgcca	1739		
DB	1320	gcgggaacacacatcgccacacatttaaggtctatcgagcgtatgcagttctggtgcca	1379		
QY	1740	gaagaaattccagcaagcggaagccttaagatgtggggagctcatgagcagttctc	1799		
DB	1380	gaagaaattccagcaagcggaagccttaagatgtggggagctcatgagcagttctc	1439		
QY	1800	gcagggccacactcaacctcatggtgcgcatcaagagctgcagagggctggaccagtc	1859		
DB	1440	gcagggccacactcaacctcatggtgcgcatcaagagctgcagagggctggaccagtc	1499		
QY	1860	cattgggaagccctcactgttcatctccgtctcagaaaagagacaggtatcgcggaagaa	1919		
DB	1500	cattgggaagccctcactgttcatctccgtctcagaaaagagacaggtatcgcggaagaa	1559		
QY	1920	cacgtatcgcccccctgaaccagtagaagaacaggtgcagcagctggaccagagct	1979		
DB	1560	cacgtatcgcccccctgaaccagtagaagaacaggtgcagcagctggaccagagct	1619		
QY	1980	ggcactcatcacgacatgcttccacagctgctcctctgcaggtggcagcagccccgg	2039		
DB	1620	ggcactcatcacgacatgcttccacagctgctcctctgcaggtggcagcagccccgg	1679		
QY	2040	cagcgccgccccccagagagggcgggccacatcacccagcctcgcgagtgggcg	2099		
DB	1680	cagcgccgccccccagagagggcgggccacatcacccagcctcgcgagtgggcg	1739		
QY	2100	ctcgtgcacacctgagcttctcctgccagcaaacacctgccacactagcagcagctgac	2159		
DB	1740	ctcgtgcacacctgagcttctcctgccagcaaacacctgccacactagcagcagctgac	1799		
QY	2160	cgtgccagaggggcccccgatgaggggtcctgagaggggatgggctgggggatggcg	2219		
DB	1800	cgtgccagaggggcccccgatgaggggtcctgagaggggatgggctgggggatggcg	1859		
QY	2220	ctgagtgcaggggggcccagagtgccccacctggccccctctctgaaagagggcacctc	2279		
DB	1860	ctgagtgcaggggggcccagagtgccccacctggccccctctctgaaagagggcacctc	1919		
QY	2280	ctaaagggccag	2339		
DB	1920	ctaaagggccag	1979		
QY	2340	tgcttgccacagctgcacttgagggtcagcaagggcagcctctctcctggcggtgagg	2399		
DB	1980	tgcttgccacagctgcacttgagggtcagcaagggcagcctctctcctggcggtgagg	2039		
QY	2400	ggccccctcctcaggtgctgagtttaccacaaagcgccccctggccccacatggtgattg	2459		
DB	2040	ggccccctcctcaggtgctgagtttaccacaaagcgccccctggccccacatggtgattg	2099		
QY	2460	acatcactggcatggtggtgggagagagagagagagagagagagagagagagagag	2519		
DB	2100	acatcactggcatggtggtgggagagagagagagagagagagagagagagagagag	2159		
QY	2520	ccaggaagtagcacaggtgagtgagggccacacactgctggccccagggggtctcctgag	2579		
DB	2160	ccaggaagtagcacaggtgagtgagggccacacactgctggccccagggggtctcctgag	2219		

XX	09-MAR-2001 (first entry)	
DT	Human colon cancer antigen nucleotide sequence SEQ ID NO:273.	
XX		
DE		
XX		
XX	Human; colon cancer; colon cancer antigen; diagnosis; detection;	
KW	Identification; cytostatic; cardioactive; neuroprotective; vulnerary;	
KW	immunomodulatory; muscular; gynaecological; gastrointestinal;	
KW	nephrotropic; antiinfective; antibacterial; gene therapy; wound;	
KW	neural disorder; immune system disorder; muscular disorder;	
KW	reproductive disorder; gastrointestinal disorder; renal disorder;	
KW	infectious disease; cardiovascular disorder; ss.	
XX		
OS	Homo sapiens.	
XX		
PN	WO200055351-A1.	
XX		
PD	21-SEP-2000.	
XX		
XX	08-MAR-2000; 2000WO-US05883.	
PF		
XX		
PR	12-MAR-1999; 99US-0124270.	
XX		
XX	(HUMA-) HUMAN GENOME SCI INC.	
PA		
XX	Rosen CA, Ruben SM;	
PI		
XX		
DR	WPI: 2000-587534/55.	
DR	P-PSDB; AAB53506.	
XX		
PT	Colon cancer associated gene sequences, referred to as colon cancer	
PT	antigens, useful for the treatment, prevention, and diagnosis of colon	
PT	disorders such as colon cancer -	
XX		
PS	Claim 1; Page 698-699; 2104pp; English.	
XX		
XX	AAC97991 to AAC98763 encode the human colon cancer associated proteins,	
CC	called human colon cancer antigens, given in AAB53334 to AAB54006. The	
CC	human colon cancer antigens can have cytostatic, cardioactive, muscular;	
CC	neuroprotective, immunomodulatory, gynaecological, gastrointestinal, and	
CC	vulnerary, nephrotropic, antiinfective and antibacterial activities, and	
CC	can be used in gene therapy. The colon cancer antigen polynucleotides,	
CC	proteins and antibodies to the proteins are useful for the prevention,	
CC	treatment and diagnosis of colon disorders, such as colon cancer. The	
CC	polynucleotides may be used in diagnostics and research, such as for	
CC	chromosome identification, and as hybridisation probes. The proteins	
CC	may also be used to prevent diseases such as neural disorders, immune	
CC	system disorders, muscular disorders, reproductive disorders,	
CC	gastrointestinal disorders, wounds, renal disorders, infectious	
CC	diseases, and cardiovascular disorders, AAC98764 to AAC98772 and	
CC	AAB54007 represent sequences used in the exemplification of the present	
CC	invention.	
XX		
XX	Sequence 432 BP; 105 A; 120 C; 109 G; 90 T; 8 other;	

```
Query Match          5.0%; Score 159; DB 21; Length 432;
Best Local Similarity 100.0%; Pred. NO. 2.8e-62;
Matches 159; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

Qy	1878	gttcattctcgtctcagaaaaagacaagatcgcggcagcaacacgcatcgcgcgcgcct	1937
Db	2	gttcattctcgtctcagaaaaagacaagatcgcggcagcaacacgcatcgcgcgcgcct	61
Qy	1938	gaaccagtagaagacaagggtgacgcagctggacagagctggcactcatcaccgacat	1997
Db	62	gaaccagtagaagacaagggtgacgcagctggacagagctggcactcatcaccgacat	121
Qy	1998	gcttcacacagctgctctccttgcagcgtggcagcacccc	2036
Db	122	gcttcacacagctgctctccttgcagcgtggcagcacccc	160

Db 122 qcttcaccagctgctctccttgccacggtggcagacccc 160

RESULT	10
AAT94004/C	
ID	AAT94004 standard; DNA; 2821 BP.
XX	
XX	
AC	AAT94004;
XX	
XX	
DT	28-FEB-1998 (first entry)
XX	
DE	DNA encoding human KVLQTL.
XX	
KW	KVLQTL; long QT syndrome; arrhythmia; minK; potassium channel;
KW	diagnosis; therapy; human; ds.

AA
OS
Homo sapiens.

	Key	Location/Qualifiers
CD5	88..1833	
FT		/*tag= a

XX PN WO9723632-A1.

03-JUL-1997.

AA	20-DEC-1996;	96WO-US19917.
PF		

AA 29-OCT-1996: 96US-0739383.

PR 22-DEC-1995; 9505-0019014.
XX

PA (GENZ) GENZYME GENETICS.
PA (UTAH) UNIV UTAH RES FOUND.

PI Connors TD, Curran ME, Keating MF, Landes GM;

AA
DR WPI: 1997-402191/37.

XX
DR P-PSDB; AAW33333.

PT New isolated human potassium channel gene, KV1QT1, - used to develop
PT products for diagnosis, prevention and therapy of long QT syndrome
XX
PS Claim 2; page 76-79; 117pp; English.

This cDNA sequence includes a full-length coding sequence for human KvLTQ1 (see AAW3355), a novel cardiac potassium channel protein. The sequence was assembled from partial clones isolated from human pancreatic and cardiac cDNA libraries. KvLTQ1 was mapped to chromosome 11p15.5 making it a candidate for the long QT syndrome (LQT) gene. LQT is an inherited cardiac arrhythmia. 15 Families with mutations in KvLTQ1 have been identified and it was shown that in all 16 families there was complete linkage between LQT1 and KvLTQ1. The KvLTQ1 gene product coassembles with human mink to form a cardiac IKs potassium channel. IKs dysfunction is a cause of cardiac arrhythmia. Coexpression of KvLTQ1 and mink in a host cell provides a means for screening for drugs useful in treating or preventing LQT. Analysis of the KvLTQ1 gene will provide an early diagnosis of subjects with LQT. A claimed method of assessing the risk in a human for LQT syndrome comprises screening for a mutation in the KvLTQ1 gene. Transgenic animals expressing human mink and KvLTQ1 can be used to test therapeutic agents against LQT.

SQ Sequence 2821 BP; 562 A; 892 C; 843 G; 524 T; 0 other;

Query Match 3.0%; Score 96; DB 18; Length 2821;
Best Local Similarity 100.0%; Pred. No. 8.3e-34;
Matches 96; Conservative 0; Mismatches 0; Indels

Qy	407	ggcgccggtgagccttagaccgcgctctccatctacagcacgcgcccggtgttg 	466
Db	121	GGCCGCCGTTAGCCTTAGACCGCGGTCTCATCTACAGCACGCGCCCGGTGTGG 	62

Qy 467 cgcgacccacgtccagggccgctctacaactcc 502

Db 61 CGGCACCCACGTCCAGGGCCGCTACAACCTCC 26

RESULT 11
AAT90730/c
ID AAT90730 standard; cDNA; 2821 BP.
XX AC
XX AAT90730;
DT 12-FEB-1998 (first entry)
XX
DE Human KVLQT1 full-length cDNA.
XX
XX KVLQT1; long QT syndrome; arrhythmia; minK; potassium channel;
KW diagnosis; therapy; human; ds.
XX
XX Homo sapiens.
OS
XX Key Location/Qualifiers
FH 88.1833
FT CDS /*tag= a
FT
XX WO9723598-A2.
PN
XX
PD 03-JUL-1997.
XX
XX 20-DEC-1996; 96WO-US19756.
XX
XX 29-OCT-1996; 96US-0730383
PR 22-DEC-1995; 95US-0019014.
XX
XX (UTAH) UNIV UTAH RES FOUND.
XX
XX Curran ME, Keating MT, Sanguinetti MC;
XX
XX WPI; 1997-402190/37.
DR P-PSDB; AAW30038.
XX
XX Human minK and Xenopus KVLQT1 coding sequences - used for assays for
PT identifying drugs which can be used for preventing or treating long
PT QT syndrome
XX
XX Example 9; Page 76-79; 105pp; English.
PS
XX This cDNA sequence includes a full-length coding sequence for human
CC KVLQT1 (see AAW30038), a novel cardiac potassium channel protein.
CC The sequence was assembled from partial clones isolated from human
CC pancreatic and cardiac cDNA libraries. KVLQT1 was mapped to
CC chromosome 11p15.5 making it a candidate for the long QT syndrome
CC (LQT) gene. LQT is an inherited cardiac arrhythmia. One intragenic
CC deletion and 10 different missense mutations which cause LQT have
CC been identified in KVLQT1. The KVLQT1 gene product coassembles
CC with human minK to form a cardiac IKs potassium channel.
CC Coexpression of these 2 proteins in a host cell provides a means
CC for screening for drugs useful in treating or preventing LQT. The
CC products can also be used for studying mechanisms underlying common
CC arrhythmias and for presymptomatic diagnosis of LQT. Transgenic
CC animals expressing human minK and KVLQT1 can be used to test
CC therapeutic agents against LQT.
XX
XX Sequence 2821 BP; 562 A; 892 C; 843 G; 524 T; 0 other;
SQ

Query Match 3.0%; Score 96; DB 18; Length 2821;
Best Local Similarity 100.0%; Pred. No. 8.3e-34;
Matches 96; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 407 ggcgcgcggtgagcctagaccgcgcgtctccatctacagcagcgcgcgcggtgttg 466
Db 121 ggcgcgcgcggtgagcctagaccgcgcgtctccatctacagcagcgcgcgcggtgttg 62
Qy 467 ggcgcgcgcggtgagcctagaccgcgcgtctccatctacagcagcgcgcgcggtgttg 502
Db 61 ggcgcgcgcggtgagcctagaccgcgcgtctccatctacagcagcgcgcgcggtgttg 26

RESULT 12
AAZ11946
ID AAZ11946 standard; DNA; 45 BP.
XX AC
XX AAZ11946;
DT 30-NOV-1999 (first entry)
XX
DE Human potassium channel pore domain DNA sequence 6.
XX
XX Potassium channel; ataxia; arrhythmia; epilepsy; Bartter's syndrome;
KW cardiovascular disorder; CNS disorder; renal disorder; ss.
XX
XX Synthetic.
OS Homo sapiens.
XX WO9943696-A1.
PN
XX 02-SEP-1999.
PD
XX 22-FEB-1999; 99WO-US03826.
XX
XX 19-JAN-1999; 99US-0116448.
PR 25-FEB-1998; 98US-0076887.
PR 07-AUG-1998; 98US-0095836.
XX
XX (AAZ11946) AXYS PHARM INC.
XX
XX Curran ME, Hu P, Miller AP, Rutter M, Wang J;
PI WPI; 1999-527591/44.
XX
XX New nucleic acids encoding mammalian K-Hnov potassium channel
PT proteins, useful for the diagnosis and treatment of episodic ataxia
PT with myokymia, cardiac arrhythmia, epilepsy and Bartter's syndrome
XX
XX Example 1; Page 31; 112pp; English.
XX
XX This sequence represents a DNA encoding a pore domain from a
CC human potassium channel and was used in the identification and
CC isolation of human K-Hnov cDNAs (AAZ11946-211915). K-Hnov proteins
CC have a high degree of homology to known potassium channels and
CC may be alpha subunits, which form the functional channel, or
CC accessory subunits that act to modulate the channel activity. K-Hnov
CC cDNAs were isolated by extension of expressed sequence tags (ESTs) which
CC were related but not identical to known human potassium channels.
CC Potential polymorphisms detected as sequence variants between multiple
CC independent clones. Potassium channels have critical roles in various
CC cell types and biochemical pathways. Defective potassium channels are
CC known to cause four human diseases: episodic ataxia with myokymia;
CC cardiac arrhythmia (long QT syndrome); epilepsy; and Bartter's syndrome.
CC As potassium channels are critical components of virtually all cells,
CC it is likely that abnormal potassium channels are also implicated in
CC certain renal, cardiovascular and central nervous system (CNS)
CC disorders. Nucleotides encoding K-Hnov proteins may be used for
CC identifying homologous or related proteins and the DNA sequences encoding
CC them. They may be used to produce compositions that modulate the
CC expression and function of the K-Hnov protein and in studying the
CC biochemical pathways associated with it. They may also be used for the
CC recombinant production of K-Hnov protein in fermentation cultures.
CC Additionally, such nucleotides may be used in gene therapy protocols for
CC the treatment of diseases associated with abnormal potassium channels.
XX
XX Sequence 45 BP; 9 A; 10 C; 18 G; 8 T; 0 other;
SQ

Query Match 1.4%; Score 45; DB 20; Length 45;
Best Local Similarity 100.0%; Pred. No. 1.3e-10;
Matches 45; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1072 tgggtgggggtggtcacagtcaccacacatcgctatggggacaag 1116

Db 1 tgggtgggggtgggtcacagtcaccacccatcggtatgggacaag 45
|||||

RESULT 13
AAT26420
ID AAT26420 standard; cDNA to mRNA; 83 BP.
XX
AC AAT26420;
XX
DT 06-DEC-1996 (first entry)
XX
DE Human gene signature HUMGS08661.
XX

XX Gene signature; messenger RNA; mRNA; relative abundance; frequency;
KW human; cloning; mapping; non-biased library; diagnosis; detection;
KW cell typing; abnormal cell function; ss.
XX
OS Homo sapiens.
XX

XX W09514772-A1.
XX
XX 01-JUN-1995.
XX
XX 11-NOV-1994; 94WO-JP01916.
XX

XX 12-NOV-1993; 93JP-0355504.
XX (MATS/) MATSUBARA K.
XX (OKUB/) OKUBO K.
XX
XX Matsubara K, Okubo K;
XX
XX WPI: 1995-206931/27.
XX

XX Identifying gene signatures in 3'-directed human cDNA library - e.g.
PT for diagnosis of abnormal cell function, by preparing cDNA that
PT reflects relative abundance of corresp. mRNA in specific human
PT tissues
XX
XX
XX

XX Claim 1; Page 2080; 2245pp; Japanese.
XX
XX A single-stranded DNA (or its complementary strand or the corresp.
CC double-stranded DNA) which comprises one of the 7837 "GS" sequences
CC given in AAT19001-T26837 and which is able to hybridise to part of
CC human genomic DNA, cDNA or mRNA is claimed. The GS (Gene Signature)
CC sequences were obtained from 3'-directed cDNA libraries prepared
CC from various human tissues; synthesis of cDNA was initiated from the
CC 3'-end of mRNA by using poly(T) as the sole primer. Since the 3'-
CC untranslated sequence is unique to a particular mRNA species, almost
CC all the 3'-oriented cDNAs hybridise with specific mRNAs. Each library
CC is constructed so as to reflect accurately the relative abundance of
CC different mRNAs in the particular tissue from which it was derived.
CC The appearance frequency of a given GS in a cDNA library can be
CC determined (esp. using primers and probes derived from the GS
CC sequences) as a means of diagnosing abnormal cell function or for
CC recognising different cell types.
XX
XX

SQ Sequence 83 BP; 25 A; 10 C; 16 G; 31 T; 1 other;

Query Match 1.3%; Score 41; DB 16; Length 83;
Best Local Similarity 100.0%; Pred. No. 8.2e-09;
Matches 41; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 3101 gatctgtgtttaatgagtttcacagtggtattgtattat 3141
|||||

Db 1 gatctgtgtttaatgagtttcacagtggtattgtattat 41

RESULT 14
AAC89984/c
ID AAC89984 standard; cDNA; 2734 BP.

XX AAC89984;
AC
XX DT 08-MAR-2001 (first entry)
XX
DE Mutant human KVLQT1 coding sequence #1.
XX
KW Human; KVLQT1; antiarrhythmic; cardiant; gene therapy;
KW cardiac potassium channel; Jervell and Lange-Nielsen Syndrome; JLN;
KW chromosome 11p15.5; long QT syndrome; ss.
OS
XX Homo sapiens.
XX
XX US6150104-A.
XX
XX 21-NOV-2000.
XX
XX 17-AUG-1998; 98US-0135021.
XX
XX 29-JUL-1998; 98US-0094477.
XX
XX 13-JUN-1997; 97US-0874655.
XX
XX (UTAH) UNIV UTAH RES FOUND.
XX
XX Keating MT, Splawski I;
XX
XX WPI: 2001-060013/07.
XX
XX P-PSDB; AAB49499.
XX
XX DNA encoding for a mutant KVLQT1 which causes Jervell and Lange-Nielsen
PT syndrome (JLN) when homozygous, useful for diagnosing long QT syndrome,
PT or diagnosing or prognosing JLN -
XX
XX Claim 2; Columns 91-96; 58pp; English.
XX
XX KVLQT1 is a cardiac potassium channel and mutations in the KVLQT1 gene
CC cause Jervell and Lange-Nielsen Syndrome (JLN). KVLQT1 maps to
CC chromosome 11p15.5. The present sequence is a mutant KVLQT1 coding
CC sequence. The mutant KVLQT1 coding sequence is useful in the diagnosis of
CC long QT syndrome and in screening humans for the presence of KVLQT1 gene
CC variants which cause JLN syndrome.
XX
XX
SQ Sequence 2734 BP; 551 A; 864 C; 809 G; 510 T; 0 other;

Query Match 1.1%; Score 34; DB 22; Length 2734;
Best Local Similarity 100.0%; Pred. No. 9.9e-06;
Matches 34; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 407 ggcgcgcgtgagcctagaccgcgcgtctccat 440
|||||
Db 34 GGC CGCGGTGAGCCTAGACCCGCGGTCTCCAT 1

RESULT 15
AAT91065
ID AAT91065 standard; DNA; 22 BP.
XX
AC AAT91065;
XX
XX
XX DT 01-MAR-1998 (first entry)
XX
XX Human KVLQT1 S2-S3 region PCR primer 1.
XX
KW KVLQT1; long QT syndrome; arrhythmia; minK; potassium channel;
KW diagnosis; therapy; human; PCR; primer; ss.
XX
XX Synthetic.
XX
XX Homo sapiens.
XX
XX W09723632-A1.
XX
XX 03-JUL-1997.

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XX PF 20-DEC-1996; 96WO-US19917.
XX PR 29-OCT-1996; 96US-0739383.
XX PR 22-DEC-1995; 95US-0019014.
XX (GENZ ) GENZYME GENETICS.
XX PA (UTAH ) UNIV UTAH RES FOUND.
XX PI Connors TD, Curran ME, Keating MF, Landes GM;
XX WPI; 1997-402191/37.
XX New isolated human potassium channel gene, KVLQT1, - used to develop
XX products for diagnosis, prevention and therapy of long QT syndrome
XX Example 12; Page 44; 117pp; English.
XX PCR primer 1 (AAT91065) and primer 2 (AAT91066) were designed to
XX amplify DNA encoding the S2-S3 region of human KVLQT1 (see AAW33355).
XX PCR primers (AAT91065-76) were used in single-strand conformation
XX analysis (SSCP) to define mutations in the human KVLQT1 gene (see
XX AAT94004) associated with long QT syndrome (LQT). An initial SSCP
XX identified an anomalous conformer in LQT-affected members of 6 large
XX families. Further SSCP analyses identified a KVLQT1 intragenic
XX deletion and 9 missense mutations associated with LQT in small
XX families and sporadic cases.
XX Sequence 22 BP; 2 A; 3 C; 9 G; 8 T; 0 other;
SQ
Query Match 0.7%; Score 22; DB 18; Length 22;
Best Local Similarity 100.0%; Pred. No. 3.6;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 640 gagatcgtcgtggtgttct 661
Db 1 gagatcgtcgtggtgttct 22

```

Search completed: November 2, 2001, 13:36:12
Job time: 5048 sec

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OM protein - protein search, using sw model

Run on: November 2, 2001, 12:02:03 ; Search time 50.72 Seconds
(without alignments)
564.701 Million cell updates/sec

Title: US-09-135-010A-113
Perfect score: 376
Sequence: 1 MNENAINSLYEAIPLPDGS.....TWKIYIRKQRNHHIMSPSP 376

Scoring table: OUTGO
Gapop 60.0 , Gapext 60.0

Searched: 219241 seqs, 76174552 residues

Word size : 4

Total number of hits satisfying chosen parameters: 138272

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : PIR_68:*
1: pir1:*
2: pir2:*
3: pir3:*
4: pir4:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	38	10.1	645	T27186	hypothetical prote
2	21	5.6	744	T34116	voltage-gated pota
3	19	5.1	393	JC5275	voltage-gated pota
4	10	2.7	528	D63214	hypothetical 60.8
5	10	2.7	528	D86099	hypothetical prote
6	8	2.1	117	B86862	hypothetical prote
7	8	2.1	246	H82553	3-demethylubiquino
8	8	2.1	280	C82490	probable potassium
9	8	2.1	565	D72222	conserved hypothet
10	7	1.9	55	S47542	pregnancy-specific
11	7	1.9	90	S24248	Ig heavy chain v r
12	7	1.9	109	S24254	Ig heavy chain v r
13	7	1.9	109	S24253	Ig heavy chain v r
14	7	1.9	110	S24250	Ig heavy chain v r
15	7	1.9	113	S24247	Ig heavy chain v r
16	7	1.9	114	S75337	hypothetical prote
17	7	1.9	116	S75339	hypothetical prote
18	7	1.9	119	T08728	NADH dehydrogenase
19	7	1.9	164	T11377	hypothetical prote
20	7	1.9	176	T23935	conserved hypothet
21	7	1.9	184	T00277	transposase - pseu
22	7	1.9	193	S37045	hypothetical prote
23	7	1.9	241	D64511	H+-transporting AT
24	7	1.9	244	I40362	nitrate ABC transp
25	7	1.9	244	F69260	B256 protein - cas
26	7	1.9	256	1 QQOMC2	recombination prot
27	7	1.9	261	1 QBBLP	hypothetical prote
28	7	1.9	261	H85637	hypothetical prote
29	7	1.9	261	E85848	hypothetical prote

30 7 1.9 268 2 D64223 probable 1-acylgly
31 7 1.9 299 2 T20605 hypothetical prote
32 7 1.9 307 2 H82743 methionyl-tRNA for
33 7 1.9 332 2 C64164 thiamin-binding pe
34 7 1.9 335 2 S44635 f22b7.7 protein -
35 7 1.9 338 2 F69437 hypothetical prote
36 7 1.9 344 2 T18019 probable site-spec
37 7 1.9 355 2 A72331 transcription regu
38 7 1.9 357 2 T27334 hypothetical prote
39 7 1.9 376 2 S57867 oncogene 1 - human
40 7 1.9 387 2 H65132 hypothetical 44.3
41 7 1.9 395 1 G69594 cytochrome P450 bi
42 7 1.9 405 2 T21188 hypothetical prote
43 7 1.9 417 2 H83708 hypothetical prote
44 7 1.9 436 2 T03702 hypothetical prote
45 7 1.9 444 2 T26229 hypothetical prote

ALIGNMENTS

RESULT 1
T27186
hypothetical protein Y54G9A.3 - Caenorhabditis elegans
C:Species: Caenorhabditis elegans
C:Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999
C:Accession: T27186
R:Smyle, R.
submitted to the EMBL Data Library, October 1998
A:Reference number: Z20324
A:Accession: T27186
A:Status: preliminary; translated from GB/EMBL/DBDJ
A:Molecule type: DNA
A:Residues: 1-645 <WIL>
A:Cross-references: EMBL:AL032648; PIDN:CAA21699.1; GSPDB:GN000020; CESP:Y54G9A.3
A:Experimental source: clone Y54G9A
C:Genetics:
A:Gene: CESP:Y54G9A.3
A:Map position: 2
A:Introns: 56/3; 100/2; 148/1; 411/2; 541/2; 575/3

Query Match 10.1%; Score 38; DB 2; Length 645;
Best Local Similarity 100.0%; Pred. No. 1.1e-30;
Matches 38; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 215 RLGSVVFIRHQLITLYIGFLGIFSSYFYVLAEKD 252
|||||
DB 227 RLGSVVFIRHQLITLYIGFLGIFSSYFYVLAEKD 264
|||||

RESULT 2
T34116
voltage-gated potassium channel klq-1 - Caenorhabditis elegans
C:Species: Caenorhabditis elegans
C:Date: 29-Oct-1999 #sequence_revision 29-Oct-1999 #text_change 18-Feb-2000
C:Accession: T34116
R:Wilcox, B.
submitted to the EMBL Data Library, December 1995
A:Description: The sequence of C. elegans cosmid C25B8.
A:Reference number: Z21479
A:Accession: T34116
A:Status: preliminary; translated from GB/EMBL/DBDJ
A:Molecule type: DNA
A:Residues: 1-744 <WIL>
A:Cross-references: EMBL:U41556; PIDN:AACT0874.1; GSPDB:GN000028; CESP:C25B8.1
A:Experimental source: strain Bristol N2; clone C25B8
C:Genetics:
A:Gene: klq-1; CESP:C25B8.1
A:Map position: X
A:Introns: 31/3; 64/1; 81/3; 131/2; 161/3; 204/1; 262/3; 304/3; 341/3; 402/2; 426/1;

iller, L.; Grotbeck, E.J.; Davis, N.W.; Lim, A.; Dimalanta, E.; Potamousis, K.; Apod
Nature 409, 529-533, 2001
A:Title: Genome sequence of enterohemorrhagic Escherichia coli O157:H7.
A:Reference number: A85480; MUID:21074935; PMID:11206551
A:Accession: G86099
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-528 <STO>
A:Cross-references: GB:AE005174; NID:g12519015; PIDN:AAG59259.1; GSPDB:GN00145; UWG:
A:Experimental source: strain O157:H7, substrain EDL933
C:Genetics:
A:Gene: yjcc

Query Match 2.7%; Score 10; DB 2; Length 528;
Best Local Similarity 100.0%; Pred. No. 0.069;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 306 FALPAGILGS 315
Db 245 FALPAGILGS 254
|||||

RESULT 6
B86862
hypothetical protein ytgE [imported] - Lactococcus lactis subsp. lactis (strain IL140
C:Species: Lactococcus lactis subsp. lactis
C:Date: 23-Mar-2001 #sequence_revision 23-Mar-2001 #text_change 23-Mar-2001
C:Accession: B86862
R:Boilotin, A.; Wincker, P.; Mauger, S.; Jaillon, O.; Malarme, K.; Weissenbach, J.; Eh
Genome Res. in press, 2001
A:Title: The complete genome sequence of the lactic acid bacterium.
A:Reference number: A86625
A:Accession: B86862
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-117 <STO>
A:Cross-references: GB:AE005176; NID:g12724933; PIDN:AAK05996.1; GSPDB:GN00146
A:Experimental source: strain IL1403
C:Genetics:
A:Gene: ytgE

Query Match 2.1%; Score 8; DB 2; Length 117;
Best Local Similarity 100.0%; Pred. No. 2.2;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 35 ELKRETLV 42
Db 66 ELKRETLV 73
|||||

RESULT 7
H82553
3-demethylubiquinone-9 3-methyltransferase XF2471 [imported] - Xylella fastidiosa (st
C:Species: Xylella fastidiosa
C:Date: 18-Aug-2000 #sequence_revision 20-Aug-2000 #text_change 02-Sep-2000
C:Accession: H82553
R:anonymous, The Xylella fastidiosa Consortium of the Organization for Nucleotide Seq
Nature 406, 151-157, 2000
A:Title: The genome sequence of the plant pathogen Xylella fastidiosa.
A:Reference number: A82515; MUID:20365717
A:Note: for a complete list of authors see reference number A59328 below
A:Accession: H82553
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-246 <SIM>
A:Cross-references: GB:AE004055; GB:AE003849; NID:g9107661; PIDN:AAF85269.1; GSPDB:GN
A:Experimental source: strain 9a5c
R:Simpson, A.J.G.; Reinach, F.C.; Arruda, P.; Abreu, F.A.; Acencio, M.; Alvarenga, R.
Briones, M.R.S.; Bueno, M.R.P.; Camargo, A.A.; Camargo, L.E.A.; Carraro, D.M.; Carr
as-Neto, E.; Docena, C.; El-Dorriy, H.; Facincani, A.P.; Ferreira, A.J.S.
submitted to GenBank, June 2000

A:Authors: Ferreira, V.C.A.; Ferro, J.A.; Fraga, J.S.; Franca, S.C.; Franco, M.C.; Frohm J.D.; Junqueira, M.L.; Kemper, E.L.; Kitajima, J.P.; Krieger, J.E.; Kuramae, E.E.; Laion Chado, M.A.; Madeira, A.M.B.N.; Madeira, H.M.F.; Marino, C.L.; Marques, M.V.; Martins, E.A.; Authors: Martins, E.M.F.; Matsukuma, A.Y.; Menck, C.F.M.; Miracca, E.C.; Miyaki, C.Y.; F.G.; Nunes, L.R.; Oliveira, M.A.; de Oliveira, M.C.; de Oliveira, R.C.; Palmieri, D.A.; Rodrigues, V.; Rosa, A.J. de; de Rosa Jr., V.E.; de Sa, R.G.; Santelli, R.V.; Sawasak A:Authors: da Silva, A.C.R.; da Silva, F.R.; da Silva, A.M.; Silva Jr., W.A.; da Silveira M.; Tshukako, M.H.; Vallada, H.; Van Sluys, M.A.; Verjovski-Almeida, S.; Vettore, A.L.; Z A:Reference number: A59328
A:Contents: annotation
C:Genetics:
A:Gene: XF2471
C:Superfamily: 3-demethylubiquinone-9 3-O-methyltransferase; bioC homology

Query Match 2.1%; Score 8; DB 2; Length 246;
Best Local Similarity 100.0%; Pred. No. 4.1;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 134 GAEYVVRLL 141
| | | | | | | |
DB 172 GAEYVVRLL 179

RESULT 8

C82490
probable potassium channel protein VCA0194 [imported] - Vibrio cholerae (strain N16961)
C:Species: Vibrio cholerae
C:Date: 18-Aug-2000 #sequence_revision 20-Aug-2000 #text_change 02-Feb-2001
C:Accession: C82490

R:Heidelberg, J.F.; Eisen, J.A.; Nelson, W.C.; Clayton, R.A.; Gwinn, M.L.; Dodson, R.J.; Chardson, D.; Ermolaeva, M.D.; Vamathevan, J.; Bass, S.; Qin, H.; Dragoi, I.; Sellers, F. I., R.R.; Mekalanos, J.J.; Venter, J.C.; Fraser, C.M.
Nature 406, 477-483, 2000

A:Title: DNA Sequence of both chromosomes of the cholera pathogen Vibrio cholerae.

A:Reference number: A82035; MUID:20406833

A:Accession: C82490

A>Status: preliminary

A:Molecule type: DNA

A:Residues: 1-280 <HEI>

A:Cross-references: GB:AE004359; GB:AE003853; NID:g9657575; PIDN:AAF96107.1; GSPDB:GN001

A:Experimental source: serogroup O1; strain N16961; biotype El Tor

C:Genetics:

A:Gene: VCA0194

A:Map position: 2

Query Match 2.1%; Score 8; DB 2; Length 280;
Best Local Similarity 100.0%; Pred. No. 4.6;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 264 SYADALWW 271
| | | | | | | |
DB 151 SYADALWW 158

RESULT 9

D72222
conserved hypothetical protein - Thermotoga maritima (strain MSB8)

C:Species: Thermotoga maritima

C:Date: 11-Jun-1999 #sequence_revision 11-Jun-1999 #text_change 21-Jul-2000

C:Accession: D72222

R:Nelson, K.E.; Clayton, R.A.; Gill, S.R.; Gwinn, M.L.; Dodson, R.J.; Haft, D.H.; Hickey Garrett, M.M.; Stewart, A.M.; Cotton, M.D.; Pratt, M.S.; Phillips, C.A.; Richardson, D.; C.M.

Nature 399, 323-329, 1999

A:Title: Evidence for lateral gene transfer between Archaea and Bacteria from genome seq

A:Reference number: A72200; MUID:99287316

A:Accession: D72222

A>Status: preliminary

A:Molecule type: DNA

A:Residues: 1-565 <ARN>

A:Cross-references: GB:AE001809; GB:AE000512; NID:g4982257; PIDN:AAD36749.1; PID:g498225

A:Experimental source: strain MSB8
C:Genetics:
A:Gene: TM1682

Query Match 2.1%; Score 8; DB 2; Length 565;
Best Local Similarity 100.0%; Pred. No. 8.4;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 349 AAENPDSA 356
| | | | | | | |
DB 133 AAENPDSA 140

RESULT 10

S47542
pregnancy-specific beta(1)-glycoprotein-11 - human

C:Species: Homo sapiens (man)

C:Date: 26-Dec-1994 #sequence_revision 26-May-1995 #text_change 17-Mar-1999

C:Accession: S47542

R:Dee, T.W.; Mettenleiter, P.A.; Mansfield, R.C.

Biochim. Biophys. Acta 1219, 195-197, 1994

A:Title: Sequence of a novel pregnancy-specific beta(1)-glycoprotein C-terminal domain

A:Reference number: S47542; MUID:94368856

A:Accession: S47542

A>Status: preliminary; translation not shown

A:Molecule type: DNA

A:Residues: 1-55 <JOE>

A:Cross-references: EMBL:L17043

A>Note: 33-X is the translation of a stop-codon; spliced according to feature informa

C:Genetics:

A:Introns: 11/2; 19/1; 33/3; 39/2; 43/1

C:Keywords: glycoprotein

Query Match 1.9%; Score 7; DB 2; Length 55;
Best Local Similarity 100.0%; Pred. No. 12;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 163 PISVIDL 169
| | | | | | | |
DB 41 PISVIDL 47

RESULT 11

S24248
Ig heavy chain V region (VH26) - human

C:Species: Homo sapiens (man)

C:Date: 19-Feb-1994 #sequence_revision 10-Nov-1995 #text_change 23-Jul-1999

C:Accession: S24248

R:Stewart, A.K.; Huang, C.; Stollar, B.D.; Schwartz, R.S.

submitted to the EMBL Data Library, June 1992

A:Description: A single VH gene predominates in the rearranged and expressed human B

A:Reference number: S24247

A:Accession: S24248

A>Status: preliminary

A:Molecule type: DNA

A:Residues: 1-90 <STF>

A:Cross-references: EMBL:X67069; NID:g38395; PIDN:CAA47454.1; PID:g38396

C:Superfamily: immunoglobulin V region; immunoglobulin homology

C:Keywords: heterotetramer; immunoglobulin

Query Match 1.9%; Score 7; DB 2; Length 90;
Best Local Similarity 100.0%; Pred. No. 19;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 151 VGVWGRLL 157
| | | | | | | |
DB 4 VGVWGRLL 10

RESULT 12

```
S24254
Ig heavy chain V region (VH26-DXP2-JH4) - human
C:Species: Homo sapiens (man)
C:Date: 19-Feb-1994 #sequence_revision 10-Nov-1995 #text_change 21-Jan-2000
C:Accession: S24254
R:Stewart, A.K.; Huang, C.; Stollar, B.D.; Schwartz, R.S.
submitted to the EMBL Data Library, June 1992
A:Description: A single VH gene predominates in the rearranged and expressed human B cell
A:Reference number: S24247
A:Accession: S24254
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-109 <STE>
A:Cross-references: EMBL:X67062
C:Superfamily: immunoglobulin V region; immunoglobulin homology
C:Keywords: heterotetramer; immunoglobulin
F:15-97/Domain: immunoglobulin homology <IMM>

Query Match          1.9%; Score 7; DB 2; Length 109;
Best Local Similarity 100.0%; Pred. No. 22;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 151 VGVWGRL 157
Db 4 VGVWGRL 10

RESULT 13
S24253
-Ig heavy chain V region (VH26-DLR4-JH6) - human
C:Species: Homo sapiens (man)
C:Date: 19-Feb-1994 #sequence_revision 10-Nov-1995 #text_change 21-Jan-2000
C:Accession: S24253
R:Stewart, A.K.; Huang, C.; Stollar, B.D.; Schwartz, R.S.
submitted to the EMBL Data Library, June 1992
A:Description: A single VH gene predominates in the rearranged and expressed human B cell
A:Reference number: S24247
A:Accession: S24253
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-109 <STE>
A:Cross-references: EMBL:X67061
C:Superfamily: immunoglobulin V region; immunoglobulin homology
C:Keywords: heterotetramer; immunoglobulin
F:12-94/Domain: immunoglobulin homology <IMM>

Query Match          1.9%; Score 7; DB 2; Length 109;
Best Local Similarity 100.0%; Pred. No. 22;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 151 VGVWGRL 157
Db 1 VGVWGRL 7

RESULT 14
S24250
-Ig heavy chain V region (VH26-DN1-JH4) - human
C:Species: Homo sapiens (man)
C:Date: 19-Feb-1994 #sequence_revision 10-Nov-1995 #text_change 21-Jan-2000
C:Accession: S24250
R:Stewart, A.K.; Huang, C.; Stollar, B.D.; Schwartz, R.S.
submitted to the EMBL Data Library, June 1992
A:Description: A single VH gene predominates in the rearranged and expressed human B cell
A:Reference number: S24247
A:Accession: S24250
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-110 <STE>
A:Cross-references: EMBL:X67071
C:Superfamily: immunoglobulin V region; immunoglobulin homology
```

```
C:Keywords: heterotetramer; immunoglobulin
F:15-97/Domain: immunoglobulin homology <IMM>
```

```
Query Match          1.9%; Score 7; DB 2; Length 110;
Best Local Similarity 100.0%; Pred. No. 22;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
QY 151 VGVWGRL 157
Db 4 VGVWGRL 10
```

```
RESULT 15
S24247
```

```
Ig heavy chain V region (VH26-DLR2-JH3) - human
C:Species: Homo sapiens (man)
C:Date: 19-Feb-1994 #sequence_revision 10-Nov-1995 #text_change 21-Jan-2000
C:Accession: S24247
R:Stewart, A.K.; Huang, C.; Stollar, B.D.; Schwartz, R.S.
submitted to the EMBL Data Library, June 1992
A:Description: A single VH gene predominates in the rearranged and expressed human B
A:Reference number: S24247
A:Accession: S24247
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-113 <STE>
A:Cross-references: EMBL:X67060; NID:g38377; PIDN:CAA47445.1; PID:g38378
C:Superfamily: immunoglobulin V region; immunoglobulin homology
C:Keywords: heterotetramer; immunoglobulin
F:15-97/Domain: immunoglobulin homology <IMM>
```

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Query Match          1.9%; Score 7; DB 2; Length 113;
Best Local Similarity 100.0%; Pred. No. 23;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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```
QY 151 VGVWGRL 157
Db 4 VGVWGRL 10
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Search completed: November 2, 2001, 12:03:05
Job time: 62 sec
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Result No.	Score	Query Match	§			DB	ID	Description
			Length	DB	ID			
-	1	3181	100.0	3181	3	US-09-135-021-1	Sequence 1, Appli	
	2	3061	96.2	3182	3	US-09-135-021-5	Sequence 5, Appli	
	3	2702	84.9	2734	3	US-09-135-021-79	Sequence 79, Appl	
C	4	34	1.1	2734	3	US-09-135-021-79	Sequence 79, Appl	
	5	20	0.6	892	4	US-09-179-558-64	Sequence 64, Appl	
	6	20	0.6	936	4	US-09-179-558-62	Sequence 62, Appl	
C	7	20	0.6	2065	2	US-08-968-751-1	Sequence 1, Appli	
	8	19	0.6	19	3	US-09-135-021-39	Sequence 39, Appl	
	9	19	0.6	19	3	US-09-135-021-40	Sequence 40, Appl	
C	10	19	0.6	7011	1	US-08-306-691B-42	Sequence 42, Appl	
	11	18	0.6	18	3	US-09-135-021-41	Sequence 41, Appl	
	12	18	0.6	18	3	US-09-135-021-72	Sequence 72, Appl	
	13	18	0.6	18	3	US-09-135-021-73	Sequence 73, Appl	
	14	18	0.6	419	1	US-08-519-777-30	Sequence 30, Appl	
	15	18	0.6	419	1	US-08-742-035-30	Sequence 30, Appl	
	16	18	0.6	419	2	US-08-777-019-30	Sequence 30, Appl	
	17	18	0.6	419	2	US-08-777-143-30	Sequence 30, Appl	
	18	18	0.6	419	3	US-08-775-414-30	Sequence 30, Appl	
	19	18	0.6	419	4	US-08-931-858B-30	Sequence 30, Appl	
	20	18	0.6	419	4	US-08-981-739-30	Sequence 30, Appl	
	21	18	0.6	426	3	US-08-775-414-88	Sequence 88, Appl	
	22	18	0.6	450	3	US-08-775-414-90	Sequence 90, Appl	
	23	18	0.6	585	1	US-08-519-777-12	Sequence 12, Appl	
	24	18	0.6	585	1	US-08-742-035-12	Sequence 12, Appl	
	25	18	0.6	585	1	US-08-777-019-12	Sequence 12, Appl	
	26	18	0.6	585	2	US-08-777-143-12	Sequence 12, Appl	
C	27	18	0.6	585	3	US-08-775-414-12	Sequence 12, Appl	


```

Db 3001 ttgccagctgtcgtgagccgcagaggaagtgcagggttcctacacaggacagggttccctctcgtg 3060
Qy 3060 ggcattacatcgcatagaaaatcaataatttgggtgatttggatctgtgttttaataatgagt 3119
|||
Db 3061 ggcattacatcgcatagaaaatcaataatttgggtgatttggatctgtgttttaataatgagt 3120
Qy 3120 ttccacgctgtgatttttgattatttaatttgcgaagcttttccctaaataaacgctggagaatca 3179
|||
Db 3121 ttccacgctgtgatttttgattatttaatttgcgaagcttttccctaaataaacgctggagaatca 3180
Qy 3180 ca 3181
||
Db 3181 ca 3182

RESULT 3
US-09-135-021-79
; Sequence 79, Application US/09135021A
; Patent No. 6150104
; GENERAL INFORMATION:
; APPLICANT: Splawski, Igor
; APPLICANT: Keating, Mark T.
; TITLE OF INVENTION: A HOMOZYGOUS MUTATION IN KVLQT1 WHICH CAUSES JERVELL
; TITLE OF INVENTION: AND LANGE-NIELSEN SYNDROME
; FILE REFERENCE: 2323-128
; CURRENT APPLICATION NUMBER: US/09/135,021A
; CURRENT FILING DATE: 1998-08-17
; EARLIER APPLICATION NUMBER: 08/874,655
; EARLIER FILING DATE: 1997-06-13
; EARLIER APPLICATION NUMBER: 60/094,477
; EARLIER FILING DATE: 1998-07-29
; NUMBER OF SEQ ID NOS: 80
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 79
; LENGTH: 2734
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (1)..(1743)
US-09-135-021-79

```

Query Match	84.9%	Score 2702;	DB 3;	Length 2734;
Best Local Similarity	100.0%;	Pred. No. 0;		
Matches 2702;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps
Qy 480	ccagggcgcgctctacaa	ttctctcgagcgtccaccgcgtcggaatgcttcggttacca	539	
Db 33	ccagggcgcgctctacaa	ttctctcgagcgtccaccgcgtcggaatgcttcggttacca	92	
Qy 540	cttcgcgcgtctccctcat	gctcgtctcgctcatcttcacgctgctgccaccatcga	599	
Db 93	cttcgcgcgtctccctcat	gctcgtctcgctcatcttcacgctgctgccaccatcga	152	
Qy 600	gcagtatgcgcgcctggccac	ggggaactctcttgatggagatcgctggtggtgtt	659	
Db 153	gcagtatgccgcctggccac	ggggaactctcttgatggagatcgctggtggtgtt	212	
Qy 660	cttcgggacggagtagct	gtggtccgcgcctctggtccgcggtcgccgcgaagtacgtggg	719	
Db 213	cttcgggacggagtagct	gtggtccgcgcctctggtccgcggtcgccgcgaagtacgtggg	272	
Qy 720	cctctggggggcggtcg	cttttgcgcgggaagccatttcacatcatcgacctcatcggt	779	
Db 273	cctctggggggcggtcg	cttttgcgcgggaagccatttcacatcatcgacctcatcggt	332	
Qy 780	cgtaggcctccacgtggt	ctcctctcggtggcctccaaaggcgaggtgtttgcacacgtcggc	839	
Db 333	cgtaggcctccacgtg	ctcctctcggtggcctccaaaggcgaggtgtttgcacacgtcggc	392	
Qy 840	catcagggggcatccgc	cttctctcgatctcgtgagtagctacacgtcgacgcgcaggagg	899	

Db	393	catagggggaatccgcttctcgtacatcctgagatgctacacgtcgcacgcgcagggagg	452
Qy	900	cactcgaggctcctgggtccgttgcttctatccacgcgcagagctgataaacacct	959
Db	453	cactcgaggctcctgggtccgttgcttctatccacgcgcagagctgataaacacct	512
Qy	960	gtacatcggtctcctgggctcatcttctcgttactttgttacctggtcgagaagga	1019
Db	513	gtacatcggtctcctgggctcatcttctcgttactttgttacctggtcgagaagga	572
Qy	1020	cgcggtgaacgaatcaagcgcgctggagtgcgcagctacacagatcgctgtgtgtggg	1079
Db	573	cgcggtgaacgaatcaagcgcgctggagtgcgcagctacacagatcgctgtgtgtggg	632
Qy	1080	gggtgtcacagtcacacacatcgctatggggacaaagtccccagacgtggtgcggaaa	1139
Db	633	gggtgtcacagtcacacacatcgctatggggacaaagtccccagacgtggtgcggaaa	692
Qy	1140	gacatcgctctcgtcttctgtgtttgcaatctcttcttcttgcgtccccagcgggat	1199
Db	693	gacatcgctctcgtcttctgtgtttgcaatctcttcttgcgtccccagcgggat	752
Qy	1200	tcttgctcgggggttgccctgaaggtgcagacagaagcagaggcagaagcacttcaacgc	1259
Db	753	tcttgctcgggggttgccctgaaggtgcagacagaagcagaggcagaagcacttcaacgc	812
Qy	1260	gcagatcccgcgcgagctcactcatcagaccgatggagggtgctatgctgcgcgagaa	1319
Db	813	gcagatcccgcgcgagctcactcatcagaccgatggagggtgctatgctgcgcgagaa	872
Qy	1320	ccccgactctccacttggaatctacatccggaagccccccggagccacactctgct	1379
Db	873	ccccgactctccacttggaatctacatccggaagccccccggagccacactctgct	932
Qy	1380	gtacccagcccaaccccaagaagtctgtgtgtaagaaaaaaagtccaagctgga	1439
Db	933	gtacccagcccaaccccaagaagtctgtgtgtaagaaaaaaagtccaagctgga	992
Qy	1440	caagacaaatgggtgactcctcgagagaagatgctacaotccccatatccagctgcga	1499
Db	993	caagacaaatgggtgactcctcgagagaagatgctacaotccccatatccagctgcga	1052
Qy	1500	ccccccagaagcggcgtgagccactctctctgtcacgcgttatgacagttctgtaag	1559
Db	1053	ccccccagaagcggcgtgagccactctctctgtcacgcgttatgacagttctgtaag	1112
Qy	1560	gaagagcccaactgctggaagtgaagatgcccatttcatgagaaccaacagcttcgc	1619
Db	1113	gaagagcccaactgctggaagtgaagatgcccatttcatgagaaccaacagcttcgc	1172
Qy	1620	cgaggacctgacctgaagggagagactctgtgacccccatccacacatctcacagct	1679
Db	1173	cgaggacctgacctgaagggagagactctgtgacccccatccacacatctcacagct	1232
Qy	1680	gcgggaacacattcgggccaccattaaagtattcgacgcgtcagttacttttggccaa	1739
Db	1233	gcgggaacacattcgggccaccattaaagtattcgacgcgtcagttacttttggccaa	1292
Qy	1740	gaagaaattccagacgcgcgaagccttaacgatgtcgggacgtcattgagcagttactc	1799
Db	1293	gaagaaattccagacgcgcgaagccttaacgatgtcgggacgtcattgagcagttactc	1352
Qy	1800	gcagggccacctcaacctcatggtgcgcatcaagagagctgcagaggagccttgacaccagtc	1859
Db	1353	gcagggccacctcaacctcatggtgcgcatcaagagagctgcagaggagccttgacaccagtc	1412
Qy	1860	catttgggaagccctcactgttcatctccgtctcagaaaaagcagaagatcgcgcgaccaa	1919
Db	1413	catttgggaagccctcactgttcatctccgtctcagaaaaagcagaagatcgcgcgaccaa	1472
Qy	1920	caagatcggcgccgcctgaacccagtagaagacaaggtgcacgcagcttgacaccagagct	1979
Db	1473	cagatcggcgccgcctgaacccagtagaagacaaggtgcacgcagcttgacaccagagct	1532

QY 1980 ggcactcatcagacatgttccaccagctgtctctcttgacaggtggcagaccccccg 2039
Db 1533 ggcactcatcagacatgttccaccagctgtctctcttgacaggtggcagaccccccg 1592
QY 2040 cagcggcgccccccagagagggcgccacatcacccagccctgcggcagtgggcg 2099
Db 1593 cagcggcgccccccagagagggcgccacatcacccagccctgcggcagtgggcg 1652
QY 2100 ctccgtgacctgagcttctctccagcaacacccctgcccacctacagcagctgac 2159
Db 1653 ctccgtgacctgagcttctctccagcaacacccctgcccacctacagcagctgac 1712
QY 2160 cgtgccagagggcccccgatgaggggtctctgagaggggatgggggatgggc 2219
Db 1713 cgtgccagagggcccccgatgaggggtctctgagaggggatgggggatgggc 1772
QY 2220 ctgagtgaagagggagcccaagagtgccccacctctctctgaagagagccacctc 2279
Db 1773 ctgagtgaagagggagcccaagagtgccccacctctctctgaagagagccacctc 1832
QY 2280 ctaaaagggccagagagagagccacctctcagagggcccccaatacccatgacccatgc 2339
Db 1833 ctaaaagggccagagagagagccacctctcagagggcccccaatacccatgacccatgc 1892
QY 2340 tctctggcacagcctgacattgggggctcagcaagggccacctctctctggccgggtggg 2399
Db 1893 tctctggcacagcctgacattgggggctcagcaagggccacctctctctggccgggtggg 1952
QY 2400 gggcccgctcaggtctgagtggttaccacagcgccctggccccccacatggatgttg 2459
Db 1953 gggcccgctcaggtctgagtggttaccacagcgccctggccccccacatggatgttg 2012
QY 2460 acatcactggcattggtgtgggacccagtgaggagggcagggcctggccatgtatgg 2519
Db 2013 acatcactggcattggtgtgggacccagtgaggagggcagggcctggccatgtatgg 2072
QY 2520 ccaggaagttagcacaggtgagtgagggccaccctctgtggccccggggggtctctctgag 2579
Db 2073 ccaggaagttagcacaggtgagtgagggccaccctctgtggccccggggggtctctctgag 2132
QY 2580 gggagacagagcaaaccttgagcccccagcctcaaatccagagccctggccagcacagga 2639
Db 2133 gggagacagagcaaaccttgagcccccagcctcaaatccagagccctggccagcacagga 2192
QY 2640 gggcaggacagccagcagctgactacagggccaccggaataaaagccagagggccatt 2699
Db 2193 gggcaggacagccagcagctgactacagggccaccggaataaaagccagagggccatt 2252
QY 2700 tggagggcctgggctggctcctcactctcaggaatgctgacccatggggcaggagact 2759
Db 2253 tggagggcctgggctggctcctcactctcaggaatgctgacccatggggcaggagact 2312
QY 2760 tggagagactgctctgagccccagcctccagagagagagagagctcaccattccccca 2819
Db 2313 tgggagagactgctctcagccccagcctccagagagagagagagctcaccattccccca 2372
QY 2820 gggcagctggttggagtgggggaagccaccctccctgggttagactgccagctcttctct 2879
Db 2373 gggcagctggttggagtgggggaagccaccctccctgggttagactgccagctcttctct 2432
QY 2880 agctggagagagccctgctctcgcgccctgagcccatgtgcgtgggggtcccccctc 2939
Db 2433 agctggagagagccctgctctcgcgccctgagcccatgtgcgtgggggtcccccctc 2492
QY 2940 caacccctcgccagctccagcagcagcagcaaacacacagagagggactgccacctcccc 2999
Db 2493 caacccctcgccagctccagcagcagcagcaaacacacagagagggactgccacctcccc 2552
QY 3000 ttgcccagctgctgagcccgagagagtgaggttctctacacagagaggggttctctctg 3059
Db 2553 ttgcccagctgctgagcccgagagagtgaggttctctacacagagaggggttctctctg 2612

QY 3060 ggcattatccgcacataaataattttgtgtgatttgatctgtgttttaagt 3119
Db 2613 ggcattatccgcacataaataattttgtgtgatttgatctgtgttttaagt 2672
QY 3120 ttccagctggtgattttgatttaattgtcaagcttttcttaataaaacgtggagaatca 3179
Db 2673 ttccagctggtgattttgatttaattgtcaagcttttcttaataaaacgtggagaatca 2732
QY 3180 ca 3181
Db 2733 ca 2734

RESULT 4
US-09-135-021-79/c
; Sequence 79, Application US/09135021A
; Patent No. 6150104
; GENERAL INFORMATION:
; APPLICANT: Splawski, Igor T.
; TITLE OF INVENTION: A HOMOZYGOUS MUTATION IN KVLQT1 WHICH CAUSES JERVELL
; FILE REFERENCE: 2323-128
; CURRENT APPLICATION NUMBER: US/09/135,021A
; EARLIER FILING DATE: 1998-08-17
; EARLIER APPLICATION NUMBER: 08/874,655
; EARLIER FILING DATE: 1997-06-13
; EARLIER APPLICATION NUMBER: 60/094,477
; EARLIER FILING DATE: 1998-07-29
; NUMBER OF SEQ ID NOS: 80
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 79
; LENGTH: 2734
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (1)..(1743)
US-09-135-021-79

Query Match 1.1%; Score 34; DB 3; Length 2734;
Best Local Similarity 100.0%; Pred. No. 2.1e-06;
Matches 34; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 407 ggcgcggctgagcctagaccgcgcgtctccat 440
Db 34 GCGCGCGGTGAGCCTAGACCCGCGCTCTCCAT 1

RESULT 5
US-09-179-558-64
; Sequence 64, Application US/09179558
; Patent No. 6180612
; GENERAL INFORMATION:
; APPLICANT: Hockensmith, Joel W.
; APPLICANT: Muthuswami, Rohini
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
; TITLE OF INVENTION: TARGETING DNA METABOLIC PROCESSES USING
; NUMBER OF SEQUENCES: 66
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: PENNIE & EDMONDS LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: NY
; COUNTRY: USA
; ZIP: 10036-2711
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq Version 2.0

;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/09/179,558
;; FILING DATE: 27-OCT-1998
;; CLASSIFICATION: 514
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: U.S. 09/060,470
;; FILING DATE: 15-APR-1998
;; PRIOR APPLICATION DATA: U.S. 60/063,898
;; FILING DATE: 31-OCT-1997
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Coruzzi, Laura A
;; REGISTRATION NUMBER: 30,742
;; REFERENCE/DOCKET NUMBER: 9426-005-999
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: (212)7909090
;; TELEX: (212)8699741
;; INFORMATION FOR SEQ ID NO: 64:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 892 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;; MOLECULE TYPE: Other
US-09-179-558-64

Query Match 0.6%; Score 20; DB 4; Length 892;
Best Local Similarity 100.0%; Pred. No. 6.2;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 974 tgggcctcatcttctctcg 993
|||||
Db 610 TGGGCTCATCTTCTCTCG 629

RESULT 6
US-09-179-558-62
;; Sequence 62, Application US/09179558
;; Patent No. 6180612
;; GENERAL INFORMATION:
;; APPLICANT: Hockensmith, Joel W.
;; APPLICANT: Muthuswami, Rohini
;; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
;; TARGETING DNA METABOLIC PROCESSES USING
;; AMINOGLYCOSIDE DERIVATIVES
;; NUMBER OF SEQUENCES: 66
;; CORRESPONDENCE ADDRESS:
;; ADDRESSEE: PENNIE & EDMONDS LLP
;; STREET: 1155 Avenue of the Americas
;; CITY: New York
;; STATE: NY
;; COUNTRY: USA
;; ZIP: 10036-2711
;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: Diskette
;; COMPUTER: IBM Compatible
;; OPERATING SYSTEM: DOS
;; SOFTWARE: FastSeq Version 2.0
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/09/179,558
;; FILING DATE: 27-OCT-1998
;; CLASSIFICATION: 514
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: U.S. 09/060,470
;; FILING DATE: 15-APR-1998
;; PRIOR APPLICATION DATA: U.S. 60/063,898
;; FILING DATE: 31-OCT-1997
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Coruzzi, Laura A
;; REGISTRATION NUMBER: 30,742

;; REFERENCE/DOCKET NUMBER: 9426-005-999
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: (212)7909090
;; TELEX: (212)8699741
;; INFORMATION FOR SEQ ID NO: 62:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 936 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: double
;; TOPOLOGY: linear
;; MOLECULE TYPE: Other
US-09-179-558-62

Query Match 0.6%; Score 20; DB 4; Length 936;
Best Local Similarity 100.0%; Pred. No. 6.2;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 974 tgggcctcatcttctctcg 993
|||||
Db 654 TGGGCTCATCTTCTCTCG 673

RESULT 7
US-08-968-751-1/c
;; Sequence 1, Application US/08968751
;; Patent No. 5948643
;; GENERAL INFORMATION:
;; APPLICANT: Rubinfeld, Bonnee
;; APPLICANT: Polakis, Paul G.
;; APPLICANT: Ligenfelter, Carol
;; APPLICANT: Vuong, Terilyn T.
;; TITLE OF INVENTION: MODULATORS OF BRCA1 ACTIVITY
;; NUMBER OF SEQUENCES: 6
;; CORRESPONDENCE ADDRESS:
;; ADDRESSEE: ONYX Pharmaceuticals, Inc.
;; STREET: 3031 Research Drive
;; CITY: Richmond
;; STATE: CA
;; COUNTRY: USA
;; ZIP: 94806
;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: Floppy disk
;; COMPUTER: IBM PC compatible
;; OPERATING SYSTEM: PC-DOS/MS-DOS
;; SOFTWARE: PatentIn Release #1.0, Version #1.30
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/08/968,751
;; FILING DATE:
;; CLASSIFICATION: 435
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Giotta, Gregory
;; REGISTRATION NUMBER: 32,028
;; REFERENCE/DOCKET NUMBER: ONYX1024 GG
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: (510) 262-8710
;; TELEX: (510) 222-9758
;; INFORMATION FOR SEQ ID NO: 1:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 2065 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: double
;; TOPOLOGY: linear
;; MOLECULE TYPE: CDNA
;; HYPOTHETICAL: NO
;; ANTI-SENSE: NO
;; FEATURE:
;; NAME/KEY: CDS
;; LOCATION: 103..1512
US-08-968-751-1

Query Match 0.6%; Score 20; DB 2; Length 2065;
Best Local Similarity 100.0%; Pred. No. 6.1;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2548 ccacacctgcttgccagg 2567
|||||
Db 1939 CCCACCTGCTTGGCCAGG 1920

RESULT 8
US-09-135-021-39
; Sequence 39, Application US/09135021A
; Patent No. 6150104
; GENERAL INFORMATION:
; APPLICANT: Splawski, Igor
; APPLICANT: Keating, Mark T.
; TITLE OF INVENTION: A HOMOZYGOUS MUTATION IN KVLQ1 WHICH CAUSES JERVELL
; FILE REFERENCE: 2323-128
; CURRENT APPLICATION NUMBER: US/09/135,021A
; CURRENT FILING DATE: 1998-08-17
; EARLIER APPLICATION NUMBER: 08/874,655
; EARLIER FILING DATE: 1997-06-13
; EARLIER APPLICATION NUMBER: 60/094,477
; EARLIER FILING DATE: 1998-07-29
; NUMBER OF SEQ ID NOS: 80
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 39
; LENGTH: 19
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-135-021-39

Query Match 0.6%; Score 19; DB 3; Length 19;
Best Local Similarity 100.0%; Pred. No. 21;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 99 ctgcgcttcgctgcagctc 117
|||||
Db 1 ctgcgcttcgctgcagctc 19

RESULT 9
US-09-135-021-40/c
; Sequence 40, Application US/09135021A
; Patent No. 6150104
; GENERAL INFORMATION:
; APPLICANT: Splawski, Igor
; APPLICANT: Keating, Mark T.
; TITLE OF INVENTION: A HOMOZYGOUS MUTATION IN KVLQ1 WHICH CAUSES JERVELL
; FILE REFERENCE: 2323-128
; CURRENT APPLICATION NUMBER: US/09/135,021A
; CURRENT FILING DATE: 1998-08-17
; EARLIER APPLICATION NUMBER: 08/874,655
; EARLIER FILING DATE: 1997-06-13
; EARLIER APPLICATION NUMBER: 60/094,477
; EARLIER FILING DATE: 1998-07-29
; NUMBER OF SEQ ID NOS: 80
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 40
; LENGTH: 19
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-135-021-40

Query Match 0.6%; Score 19; DB 3; Length 19;
Best Local Similarity 100.0%; Pred. No. 21;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 414 ggtgagcctagaccgcgc 432

Db 19 GGTGAGCCTAGACCCGCG 1
|||||

RESULT 10
US-08-306-691B-42/c
; Sequence 42, Application US/08306691B
; Patent No. 5734039
; GENERAL INFORMATION:
; APPLICANT: Calabretta, Bruno
; APPLICANT: Skorski, Tomasz
; TITLE OF INVENTION: ANTISENSE
; TITLE OF INVENTION: OLIGONUCLEOTIDES TARGETING COOPERATING ONCOGENES
; NUMBER OF SEQUENCES: 55
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Seidel, Gonda, Lavoragna & Monaco, P.C.
; STREET: Two Penn Center, Suite 1800
; CITY: Philadelphia
; STATE: Pennsylvania
; COUNTRY: U.S.A.
; ZIP: 19102
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.50 inch, 720 Kb
; COMPUTER: IBM PS/2
; OPERATING SYSTEM: MS-DOS
; SOFTWARE: WordPerfect 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/306,691B
; FILING DATE: September 15, 1994
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Monaco, Daniel A.
; REGISTRATION NUMBER: 30,480
; REFERENCE/DOCKET NUMBER: 8321-8
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (215) 568-8383
; TELEFAX: (215) 568-5549
; TELEX: No. 5734039e
; INFORMATION FOR SEQ ID NO: 42:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7011 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
US-08-306-691B-42

Query Match 0.6%; Score 19; DB 1; Length 7011;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2346 gcacagcctgcacttgggg 2364
|||||
Db 3821 GCACAGCCTGCACCTGGGG 3803

RESULT 11
US-09-135-021-41
; Sequence 41, Application US/09135021A
; Patent No. 6150104
; GENERAL INFORMATION:
; APPLICANT: Splawski, Igor
; APPLICANT: Keating, Mark T.
; TITLE OF INVENTION: A HOMOZYGOUS MUTATION IN KVLQ1 WHICH CAUSES JERVELL
; FILE REFERENCE: 2323-128
; CURRENT APPLICATION NUMBER: US/09/135,021A
; CURRENT FILING DATE: 1998-08-17
; EARLIER APPLICATION NUMBER: 08/874,655
; EARLIER FILING DATE: 1997-06-13

; EARLIER APPLICATION NUMBER: 60/094,477
; EARLIER FILING DATE: 1998-07-29
; NUMBER OF SEQ ID NOS: 80
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 41
; LENGTH: 18
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-135-021-41

Query Match 0.6%; Score 18; DB 3; Length 18;
Best Local Similarity 100.0%; Pred. No. 60;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 372 cgcgcgccccccagttgc 389
| | | | | | | | | | | | | | | | | |
Db 1 cgcgcgccccccagttgc 18

RESULT 12
US-09-135-021-72/c
; Sequence 72, Application US/09135021A
; Patent No. 6150104
; GENERAL INFORMATION:
; APPLICANT: Splawski, Igor
; TITLE OF INVENTION: A HOMOLOGOUS MUTATION IN KVLQT1 WHICH CAUSES JERVELL
; FILE REFERENCE: 2323-128
; CURRENT APPLICATION NUMBER: US/09/135,021A
; CURRENT FILING DATE: 1998-08-17
; EARLIER APPLICATION NUMBER: 08/874,655
; EARLIER FILING DATE: 1997-06-13
; EARLIER APPLICATION NUMBER: 60/094,477
; EARLIER FILING DATE: 1998-07-29
; NUMBER OF SEQ ID NOS: 80
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 72
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-135-021-72

Query Match 0.6%; Score 18; DB 3; Length 18;
Best Local Similarity 100.0%; Pred. No. 60;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2200 gatggggctgggggatgg 2217
| | | | | | | | | | | | | | | | | |
Db 18 GATGGGGCTGGGGGATGG 1

RESULT 13
US-09-135-021-73
; Sequence 73, Application US/09135021A
; Patent No. 6150104
; GENERAL INFORMATION:
; APPLICANT: Splawski, Igor
; TITLE OF INVENTION: A HOMOLOGOUS MUTATION IN KVLQT1 WHICH CAUSES JERVELL
; FILE REFERENCE: 2323-128
; CURRENT APPLICATION NUMBER: US/09/135,021A
; CURRENT FILING DATE: 1998-08-17
; EARLIER APPLICATION NUMBER: 08/874,655
; EARLIER FILING DATE: 1997-06-13
; EARLIER APPLICATION NUMBER: 60/094,477
; EARLIER FILING DATE: 1998-07-29
; NUMBER OF SEQ ID NOS: 80
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 73

; LENGTH: 18
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (1)..(18)
US-09-135-021-73

Query Match 0.6%; Score 18; DB 3; Length 18;
Best Local Similarity 100.0%; Pred. No. 60;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 721 ctctggggggcggtgcgc 738
| | | | | | | | | | | | | | | | | |
Db 1 ctctggggggcggtgcgc 18

RESULT 14
US-08-519-777-30/c
; Sequence 30, Application US/08519777
; Patent No. 5739307
; GENERAL INFORMATION:
; APPLICANT: JOHNSON JR., EUGENE M.
; APPLICANT: MILBRANDT, JEFFREY D.
; APPLICANT: KOTZBAUER, PAUL T.
; APPLICANT: LAMPE, PATRICIA A.
; TITLE OF INVENTION: NEURTURIN AND RELATED GROWTH FACTORS
; NUMBER OF SEQUENCES: 78
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: ROGERS, HOWELL & HAFERKAMP, L.C.
; STREET: 7733 FORSYTH BOULEVARD, SUITE 1400
; CITY: ST. LOUIS
; STATE: MISSOURI
; COUNTRY: US
; ZIP: 63105-1817
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/519,777
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: HOLLAND, DONALD R.
; REGISTRATION NUMBER: 35,197
; REFERENCE/DOCKET NUMBER: 953095
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314) 727-5188
; TELEFAX: (314) 727-6092
; INFORMATION FOR SEQ ID NO: 30:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 419 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
US-08-519-777-30

Query Match 0.6%; Score 18; DB 1; Length 419;
Best Local Similarity 100.0%; Pred. No. 54;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 359 ccccgccgcgcgcgcgcgc 376
| | | | | | | | | | | | | | | | | |
Db 122 CCCC GCCGCCGCCGCCGCCG 105

RESULT 15
US-08-742-035-30/c

Search completed: November 2, 2001, 13:30:48
Job time: 4759 sec

PS Disclosure; Fig 2; 30pp; German.

XX This invention describes a novel eukaryotic expression vector (A) comprising a nucleic acid sequence (i), encoding a potassium channel subunit (ii), arranged so that it can be functionally expressed in eukaryotes. (A) are used to express bacterial potassium channels in eukaryotes, specifically for screening compounds for their ability to open, close or (in)activate the channels or to alter their biophysical properties, especially to identify potential antibiotics.

XX Sequence 21 AA;

Query Match 5.6%; Score 21; DB 21; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.5e-13;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 266 ADALWGWGVTVTTIGYDKVP 286

Db 1 adalwgvvtvttigydkvp 21

RESULT 14

AA49495
ID AAB49495 standard; Protein; 283 AA.

XX AAB49495;

XX 08-MAR-2001 (first entry)

XX Mutant human KVLQT1 #2.

XX Human; KVLQT1; antiarrhythmic; cardiant; gene therapy;
KW cardiac potassium channel; Jervell and Lange-Nielsen Syndrome; JLN;
KW chromosome 11p15.5; long QT syndrome.

XX Homo sapiens.

XX US6150104-A.

XX 21-NOV-2000.

XX 17-AUG-1998; 98US-0135021.

XX 29-JUL-1998; 98US-0094477.

XX 13-JUN-1997; 97US-0874655.

XX (UTAH)-UNIV UTAH RES FOUND.

XX Keating-MT, Splawski I;

XX WPI; 2001-060013/07.

XX N-PSDB; AAC89914.

XX DNA encoding for a mutant KVLQT1 which causes Jervell and Lange-Nielsen syndrome (JLN) when homozygous, useful for diagnosing long QT syndrome, or diagnosing or prognosing JLN -

XX Example 4; Columns 67-70; 58pp; English.

XX KVLQT1 is a cardiac potassium channel and mutations in the KVLQT1 gene cause Jervell and Lange-Nielsen Syndrome (JLN). KVLQT1 maps to chromosome 11p15.5. The present sequence is a mutant KVLQT1. The coding sequence for the present protein is useful in the diagnosis of long QT syndrome and in screening humans for the presence of KVLQT1 gene variants which cause JLN syndrome.

XX Sequence 283 AA;

Query Match 5.6%; Score 21; DB 22; Length 283;
Best Local Similarity 100.0%; Pred. No. 2.4e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 73 QGRVYNFLRPTGKCFVYHF 93

Db 107 qgrvynflrptgwkcfvyhf 127

RESULT 15

AA498342
ID AAY08342 standard; Protein; 807 AA.

XX AAY08342;

XX 22-JUL-1999 (first entry)

XX Human nKTQ1 protein.

XX KCNQ2; KCNQ3; human; murine; potassium channel; diagnosis; prognosis;
KW benign familial neonatal epilepsy; BFNE; juvenile myotonic epilepsy;
KW JME; rolandic epilepsy; mutant; treatment; screening; epilepsy;
KW detection; gene therapy; drug screening; nKTQ1.

XX Homo sapiens.

XX W09921875-A1.

XX 06-MAY-1999.

XX 23-OCT-1998; 98WO-US22375.

XX 24-OCT-1997; 97US-0063147.

XX (UTAH) UNIV UTAH RES FOUND.

XX Charlier C, Leppert MF, Singh NA;

XX WPI; 1999-312938/26.

XX Nucleic acid encoding potassium channels KCNQ2 and 3

XX Disclosure; Page 125-128; 195pp; English.

XX This invention describes novel human and mouse potassium channel proteins KCNQ2 and KCNQ3. Detecting mutations in sequences that encode KCNQ2 or KCNQ3, or the loss of one copy of these genes, is used for diagnosis and prognosis of benign familial neonatal epilepsy (BFNE), juvenile myotonic epilepsy (JME) or rolandic epilepsy (RE). Cells (or transgenic animals) that express wild-type or mutant KCNQ2 or 3 (also the proteins themselves in cell-free form) are used to screen for agents that can be used to treat or prevent these forms of epilepsy. Fragments of the encoding nucleic acids are used as probes or primers, either for detecting mutations or for isolation of related sequences, while the complete sequences may be used in gene therapy to provide wild-type protein. Antibodies specific for mutant or wild-type proteins are used as diagnostic reagents and for drug screening. The KCNQ2 and 3 proteins are useful in rational design of drugs and therapeutically (in replacement therapies). The forms of epilepsy associated with mutations in KCNQ2 and 3 sequences can now be diagnosed early (before symptoms are manifest), and better treatment options will be available.

XX Sequence 807 AA;

Query Match 5.6%; Score 21; DB 20; Length 807;
Best Local Similarity 100.0%; Pred. No. 5.9e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 303 ISFFALPAGILSGFALKVQQ 323

Db 333 isffalpagilsgfalkvqq 353

Search completed: November 3, 2001, 13:19:10

XX 21-NOV-2000.
 XX 17-AUG-1998; 98US-0135021.
 PF 29-JUL-1998; 98US-0094477.
 XX 13-JUN-1997; 97US-0874655.
 XX (UTAH) UNIV UTAH RES FOUND.
 XX Keating-WT, Splawski I;
 PI WPI; 2001-060013/07.
 DR N-PSDB; AAC89911.
 XX DNA encoding for a mutant KVLQT1 which causes Jervell and Lange-Nielsen
 PT syndrome (JLN) when homozygous, useful for diagnosing long QT syndrome,
 PT or diagnosing or prognosing JLN -
 XX Example 4; Columns 59-64; 58pp; English.
 XX The present sequence is wild-type human KVLQT1. KVLQT1 is a cardiac
 CC potassium channel and mutations in the KVLQT1 gene cause Jervell and
 CC Lange-Nielsen Syndrome (JLN). KVLQT1 maps to chromosome 11p15.5. The
 CC present invention relates to a mutant KVLQT1 coding sequence (see
 CC AAC89914). The mutant KVLQT1 coding sequence is useful in the diagnosis
 CC of long QT syndrome and in screening humans for the presence of KVLQT1
 CC gene variants which cause JLN syndrome.
 XX Sequence 676 AA;
 SQ

Query Match 18.4%; Score 69; DB 22; Length 676;
 Best Local Similarity 100.0%; Pred. No. 7.1e-59;
 Matches 69; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 185 GQVFATSAIRGIRFLQILRLMLHVDROGGTWRLLGSGVFIHRQELITLTYIGFLGLIFSSY 244
 Db 219 gqvfatssairgfrilqlrlmlhvdrggtwrlgsvvfihrqelittlyigflglifssy 278
 QY 245 FVYLAEKDA 253
 Db 279 fvylaekda 287

RESULT 12
 ID AAY08343 standard; Protein; 677 AA.
 XX
 AC AAY08343;
 XX
 DT 22-JUL-1999 (first entry)
 XX
 DE Human KCNQ1 protein.
 XX
 KW KCNQ2; KCNQ3; human; murine; potassium channel; diagnosis; prognosis;
 KW benign familial neonatal epilepsy; BFNE; juvenile myotonic epilepsy;
 KW JME; rolandic epilepsy; mutant; treatment; screening; epilepsy;
 KW detection; gene therapy; drug screening; KCNQ1.
 XX
 OS Homo sapiens.
 XX
 PN W09521875-A1.
 XX
 PD 06-MAY-1999.
 XX
 PF 23-OCT-1998; 98WO-US22375.
 XX
 PR 24-OCT-1997; 97US-0063147.
 XX
 PA (UTAH) UNIV UTAH RES FOUND.
 XX
 PI Charlier C, Leppert MF, Singh NA;

XX WPI; 1999-312938/26.
 XX Nucleic acid encoding potassium channels KCNQ2 and 3
 XX Disclosure; Page 128-130; 195pp; English.
 XX This invention describes novel human and mouse potassium channel proteins
 CC KCNQ2 and KCNQ3. Detecting mutations in sequences that encode KCNQ2 or
 CC KCNQ3, or the loss of one copy of these genes, is used for diagnosis and
 CC prognosis of benign familial neonatal epilepsy (BFNE), juvenile myotonic
 CC epilepsy (JME) or rolandic epilepsy (RE). Cells (or transgenic animals)
 CC that express wild-type or mutant KCNQ2 or 3 (also the proteins themselves
 CC in cell-free form) are used to screen for agents that can be used to
 CC treat or prevent these forms of epilepsy. Fragments of the encoding
 CC nucleic acids are used as probes or primers, either for detecting
 CC mutations or for isolation of related sequences, while the complete
 CC sequences may be used in gene therapy to provide wild-type protein.
 CC Antibodies specific for mutant or wild-type proteins are used as
 CC diagnostic reagents and for drug screening. The KCNQ2 and 3 proteins are
 CC useful in rational design of drugs and therapeutically (in replacement
 CC therapies). The forms of epilepsy associated with mutations in KCNQ2 and
 CC 3 sequences can now be diagnosed early (before symptoms are manifest),
 CC and better treatment options will be available.
 XX Sequence 677 AA;
 SQ

Query Match 18.4%; Score 69; DB 20; Length 677;
 Best Local Similarity 100.0%; Pred. No. 7.1e-59;
 Matches 69; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 185 GQVFATSAIRGIRFLQILRLMLHVDROGGTWRLLGSGVFIHRQELITLTYIGFLGLIFSSY 244
 Db 220 gqvfatssairgfrilqlrlmlhvdrggtwrlgsvvfihrqelittlyigflglifssy 279
 QY 245 FVYLAEKDA 253
 Db 280 fvylaekda 288

RESULT 13
 ID AAB11385 standard; Protein; 21 AA.
 XX
 AC AAB11385;
 XX
 DT 22-FEB-2001 (first entry)
 XX
 DE Potassium channel protein KVLQT1 P region.
 XX
 KW L¹T B; potassium channel protein; screening; antibiotic.
 XX
 OS Unidentified.
 XX
 PN EP1046708-A1.
 XX
 PD 23-OCT-2000.
 XX
 PF 13-APR-2000; 2000EP-0107916.
 XX
 PR 23-APR-1999; 99DE-1020044.
 XX
 PA (GENI-) FORSCHUNGSESELLSCHAFT GENION MBH.
 XX
 PI Pongs O;
 XX
 DR WPI; 2000-657763/64.
 XX
 PT Expression vector for bacterial potassium channel that is functional in
 PT eukaryotic cells, used to screen for channel modulators and potential
 PT antibiotics -
 XX

XX Human KVLQT1 associated with long QT syndrome.
 DE KVLQT1; long QT syndrome; arrhythmia; mink; potassium channel;
 KW diagnosis; therapy; human.
 XX Homo sapiens.

XX Key Location/Qualifiers
 FT 28..49
 FT /label= S1
 FT /note= "transmembrane domain"
 FT 53..75
 FT /label= S2
 FT /note= "transmembrane domain"
 FT 103..121
 FT /label= S3
 FT /note= "transmembrane domain"
 FT 126..144
 FT /label= S4
 FT /note= "transmembrane domain"
 FT 168..187
 FT /label= S5
 FT /note= "transmembrane domain"
 FT 194
 FT /note= "N-glycosylated"
 FT 206..225
 FT /label= Pore
 FT 230..259
 FT /label= S6
 FT /note= "transmembrane domain"

XX WO9723598-A2.
 XX 03-JUL-1997.
 XX 20-DEC-1996; 96WO-US19756.
 XX 29-OCT-1996; 96US-0739383.
 XX 22-DEC-1995; 95US-0019014.
 XX (UTAH) UNIV UTAH-RES. FOUND.
 XX Curran ME, Keating MT, Sanguinetti MC;
 WPI; 1997-402190/37.

XX Human mink and Xenopus KVLQT1 coding sequences - used for assays for
 PT identifying drugs which can be used for preventing or treating long
 PT QT syndrome

XX Example 8; Fig 3A; 105pp; English.
 XX This protein comprises a novel human cardiac potassium channel
 CC protein. It is encoded by the KVLQT1 gene (see AAG90730) that
 CC is associated with long QT syndrome (LQT) gene, an inherited
 CC cardiac arrhythmia. KVLQT1 protein coassembles with human mink
 CC to form the cardiac IKs potassium channel. IKs dysfunction is
 CC a cause of cardiac arrhythmia. Coexpression of KVLQT1 and mink
 CC in a host cell provides a means for screening for drugs useful in
 CC treating or preventing LQT. The products can also be used for
 CC studying mechanisms underlying common arrhythmias and for
 CC presymptomatic diagnosis of LQT. Transgenic animals that express
 CC human mink and KVLQT1 can be used to test therapeutic agents
 CC against LQT.

XX Sequence 581 AA;
 XX
 XX Query Match 18.4%; Score 69; DB 18; Length 581;
 XX Best Local Similarity 100.0%; Pred. No. 6.2e-59;
 XX Matches 69; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 185 GQVFATSAIRGIRFLOIRMLHVDROGQTRLLGWSVFIHQELITTLTYIGFLGLIFSSY 244
 DB 124 gqvfatasaigrirfqlrlmhlvdrgqgtrllgswvfihrqelittlyigflglifssy 183
 QY 245 FVYLAEKDA 253
 DB 184 fvylaekda 192

RESULT 8
 AAB49499
 ID AAB49499 standard; Protein; 581 AA.
 XX
 AC AAB49499;
 XX
 DT 08-MAR-2001 (first entry)
 XX
 DE Mutant human KVLQT1 #1.
 XX
 KW Human; KVLQT1; antiarrhythmic; cardiant; gene therapy;
 KW cardiac potassium channel; Jervell and Lange-Nielsen Syndrome; JLN;
 KW chromosome 11p15.5; long QT syndrome.
 XX
 OS Homo sapiens.
 XX
 PN US6150104-A.
 XX
 PD 21-NOV-2000.
 XX
 PF 17-AUG-1998; 98US-0135021.
 XX
 PR 29-JUL-1998; 98US-0094477.
 PR 13-JUN-1997; 97US-0874655.
 XX
 XX (UTAH) UNIV UTAH RES FOUND.
 XX
 PI Keating MT, Splawski I;
 XX
 DR WPI; 2001-060013/07.
 DR N-PSDB; AAC89984.
 XX
 PT DNA encoding for a mutant KVLQT1 which causes Jervell and Lange-Nielsen
 PT syndrome (JLN) when homozygous, useful for diagnosing long QT syndrome,
 PT or diagnosing or prognosing JLN -
 XX
 PS Claim 1; Columns 95-100; 58pp; English.
 XX
 CC KVLQT1 is a cardiac potassium channel and mutations in the KVLQT1 gene
 CC cause Jervell and Lange-Nielsen Syndrome (JLN). KVLQT1 maps to
 CC chromosome 11p15.5. The present sequence is a mutant KVLQT1. The coding
 CC sequence for the present protein is useful in the diagnosis of long QT
 CC syndrome and in screening humans for the presence of KVLQT1 gene variants
 CC which cause JLN syndrome.

XX
 SQ Sequence 581 AA;
 XX
 XX Query Match 18.4%; Score 69; DB 22; Length 581;
 XX Best Local Similarity 100.0%; Pred. No. 6.2e-59;
 XX Matches 69; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 185 GQVFATSAIRGIRFLOIRMLHVDROGQTRLLGWSVFIHQELITTLTYIGFLGLIFSSY 244
 DB 124 gqvfatasaigrirfqlrlmhlvdrgqgtrllgswvfihrqelittlyigflglifssy 183
 QY 245 FVYLAEKDA 253
 DB 184 fvylaekda 192
 RESULT 9
 AAY57368
 ID AAY57368 standard; Protein; 676 AA.

OS Homo sapiens.
 XX WO200006600-A1.
 PN 10-FEB-2000.
 PD 06-OCT-1998; 98WO-US17838.
 XX 29-JUL-1998; 98US-0094477.
 PR 17-AUG-1998; 98US-0135020.
 XX (UTAH) UNIV UTAH RES FOUND.
 PA Keating MT, Sanguinetti MC, Splawski I;
 PI WPI; 2000-195262/17.
 XX
 DR Mutant forms of genes encoding mink protein and KVLQT1 protein involved
 XX in cardiac potassium channel formation useful for screening drugs, for
 XX preventing and treating cardiac arrhythmia
 XX
 XX Disclosure; Fig 10; 167pp; English.
 XX
 XX The invention relates to KVLQT1 and KCNE1 genes, associated with long
 XX QT (LQT) syndrome. It provides a mink protein comprising a mutation which
 XX substitutes the wild type amino acids with leu, Asp, leu, His, Trp and
 XX Ala or Thr at residues 74,76,28,32,98 and 127 respectively. Screening
 XX KVLQT1 and KCNE1 is useful for identifying mutations for diagnosing and
 XX treating LQT. The ability to predict LQT enables physicians to prevent
 XX the diseases with medical therapy such as beta blocking agents and opts
 XX for better treatments. The present sequence represents the human
 XX KVLQT1 protein fragment.
 XX
 XX Sequence 570 AA:
 SQ
 Query Match 18.4%; Score 69; DB 21; Length 570;
 Best Local Similarity 100.0%; Pred. No. 6.1e-59;
 Matches 69; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 185 GQVFATSAIRGIRFQILRLMLHVDROGGTWRLLGSVVFHRRQELITTLTYIGFLGLIFSSY 244
 DB 113 gqvfatssairgfrqlrlmlhvdrggtwrlgsvvfhrqelittlyigflglifssy 172
 QY 245 FVYLAEKDA 253
 DB 173 fvyllaekda 181
 RESULT 6
 AAW33355
 ID AAW33355 standard; Protein; 581 AA.
 XX
 XX AAW33355;
 XX 28-FEB-1998 (first entry)
 XX Human KVLQT1 associated with long QT syndrome.
 XX KVLQT1; long QT syndrome; arrhythmia; mink; potassium channel;
 KW diagnosis; therapy; human.
 XX Homo sapiens.
 XX
 XX Key Location/Qualifiers
 FT Domain 28...49
 FT /label= S1
 FT /note= "transmembrane domain"
 FT Domain 53..75
 FT /label= S2
 FT /note= "transmembrane domain"
 FT Domain 103..121
 FT /label= S3

FT Domain /note= "transmembrane domain"
 FT 126..144
 FT /label= S4
 FT /note= "transmembrane domain"
 FT 168..187
 FT /label= S5
 FT /note= "transmembrane domain"
 FT Misc-difference 194
 FT /note= "N-glycosylated"
 FT 206..225
 FT /note= "pore domain"
 FT 230..259
 FT /label= S6
 FT /note= "transmembrane domain"
 XX
 XX WO9723632-A1.
 XX
 XX 03-JUL-1997.
 PD
 XX 20-DEC-1996; 96WO-US19917.
 PF
 XX 29-OCT-1996; 96US-0739383.
 PR
 XX 22-DEC-1995; 95US-0019014.
 PR
 XX (GENZ) GENZYME GENETICS.
 PA (UTAH) UNIV UTAH RES FOUND.
 XX
 XX Connors TD, Curran ME, Keating MF, Landes GM;
 PI WPI; 1997-402191/37.
 DR N-PSDB; AAT94004.
 XX
 XX New isolated human potassium channel gene, KVLQT1, - used to develop
 XX products for diagnosis, prevention and therapy of long QT syndrome
 XX
 XX Claim 1; Page 76-78; 105pp; English.
 XX
 XX This protein comprises a novel human cardiac potassium channel
 XX protein. It is encoded by the KVLQT1 gene (see AAT94004) that
 XX is associated with long QT syndrome (LQT) gene, an inherited
 XX cardiac arrhythmia. KVLQT1 protein coassembles with human mink
 XX to form the cardiac IKs potassium channel. IKs dysfunction is
 XX a cause of cardiac arrhythmia. Coexpression of KVLQT1 and mink
 XX in a host cell provides a means for screening for drugs useful in
 XX treating or preventing LQT. The products can also be used for
 XX studying mechanisms underlying common arrhythmias and for
 XX presymptomatic diagnosis of LQT. Transgenic animals that express
 XX human mink and KVLQT1 can be used to test therapeutic agents
 XX against LQT.
 XX
 XX Sequence 581 AA:
 SQ
 Query Match 18.4%; Score 69; DB 18; Length 581;
 Best Local Similarity 100.0%; Pred. No. 6.2e-59;
 Matches 69; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 185 GQVFATSAIRGIRFQILRLMLHVDROGGTWRLLGSVVFHRRQELITTLTYIGFLGLIFSSY 244
 DB 124 gqvfatssairgfrqlrlmlhvdrggtwrlgsvvfhrqelittlyigflglifssy 183
 QY 245 FVYLAEKDA 253
 DB 184 fvyllaekda 192
 RESULT 7
 AAW30038
 ID AAW30038 standard; Protein; 581 AA.
 XX
 XX AAW30038;
 XX 12-FEB-1998 (first entry)

XX 12-MAY-1999; 99WO-US10360.
 PF XX
 XX 29-JUL-1998; 98US-0094477.
 PR 17-AUG-1998; 98US-0135010.
 XX
 XX (UTAH) UNIV UTAH RES FOUND.
 PA (GENZ) GENZYME CORP.
 XX Keating-MT, Sanguinetti MC, Curran ME, Landes GM, Connors TD;
 PI Burn-TC, Splawski I;
 XX
 XX WPI; 2000-195199/17.
 DR
 XX New isolated mutant KVLQT1 nucleic acids, useful for developing
 PT products for the diagnosis, prevention and treatment of long QT
 PT syndrome -
 XX
 XX Claim 60; Fig 10; 178pp; English.
 XX
 CC The invention relates to KVLQT1 nucleic acids which have a mutation
 CC compared to wild-type KVLQT1 (AAZ8901). The KVLQT1 gene encodes a
 CC protein of 676 amino acids which forms a cardiac I(Ks) potassium channel
 CC with the KCNE1 protein (AAH80563). The KCNE1 protein has been shown to
 CC be functional in Xenopus leavis oocyte when KCNE1 DNA is injected into
 CC the egg, indicating that a homologue of the human KVLQT1 gene is present
 CC in Xenopus. The human KVLQT1 gene was then used to probe a DNA library
 CC to isolate the sequence encoding this protein. Mutations in the KVLQT1
 CC or KCNE1 genes result in cardiac arrhythmias observed as a prolonged QT
 CC curve in electrocardiograms (Long QT syndrome). The genes and proteins
 CC can be used for the diagnosis of subjects with long QT syndrome. They
 CC can also be used to screen for drugs which can be used for treating or
 CC preventing long QT syndrome. The KVLQT1 nucleic acids can be used for
 CC gene therapy, and KVLQT1 peptides can be used for peptide therapy.
 XX
 XX Sequence 376 AA;
 SQ
 Query Match 100.0%; Score 376; DB 21; Length 376;
 Best Local Similarity 100.0%; Pred. No. 0;
 Matches 376; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 MNEAINSLYEALPLPDGSSNGQROEDQANSFELKRETLVATDPRPTINLDPRVSIY 60
 DB 1 mnenainslyealplpdgssngqrqgdrgansfelkretlvatdprptinldprvsiy 60
 QY 61 SRRPLFSRTNIQGRVYNFLERTGKCFYHFTVFLVILICLIFSVLSTIQYNNLATE 120
 DB 61 srrplfsrtniqgrvynflertgkcfvyhftvflvliclifsylvstiqygnlate 120
 QY 121 TLFWMELVLFVFFGAEYVVRMLWSAGCRSKYGVGVWGRLEFARKPISVIDLIVVASVIVLC 180
 DB 121 tlfwmeilvlfvffgaeYVVRMLWSAGCRSKYGVGVWGRLEFARKPISVIDLIVVASVIVLC 180
 QY 181 VGSNGQVFATSAIRGIRFLOILRMLHVDROGQWRLGSGVVFHROELITTLTYIGFLGLI 240
 DB 181 vgsngqvfatSaIRGIRfLQILrmlhvdrggqtwrllsgsvvfhrqelittlyigflgli 240
 QY 241 FSSYFVYLAEKDAIDSSGEYQFCSYADALWGVVTTTIGYDKVQPTWIGKTIASCFSV 300
 DB 241 fssyfvyLaEKDAIDSSGEYqfcsyadAlwgvvtvtTtigydkvpqtWigktiascfsv 300
 QY 301 FAISFFALPAGILGSGFALKVQOKROKHFNRQIPAAASLIOTAWRCYAAENPDSATWKI 360
 DB 301 faisffalpagilgsgfalkvqokrqkhnfrqipaaasliqtawrcyaaenpdsatwki 360
 QY 361 YIRKQSRNHHIMSPSP 376
 DB 361 yirkqsrnhhimspsp 376
 RESULT 4
 AAY57372

ID AAY57372 standard; Protein; 137 AA.
 XX
 AC AAY57372;
 XX
 DT 19-JUN-2000 (first entry)
 XX
 DE Human KVLQT1 protein fragment.
 XX
 KW KVLQT1; KCNE1; long QT syndrome; LQT syndrome; minK protein;
 KW antiarrhythmic; gene therapy; human.
 XX
 OS Homo sapiens.
 XX
 PN WO200006600-A1.
 XX
 PD 10-FEB-2000.
 XX
 PF 06-OCT-1998; 98WO-US17838.
 XX
 PR 29-JUL-1998; 98US-0094477.
 PR 17-AUG-1998; 98US-0135020.
 XX
 PA (UTAH) UNIV UTAH RES FOUND.
 XX
 PI Keating MT, Sanguinetti MC, Splawski I;
 XX
 DR WPI; 2000-195262/17.
 XX
 XX Mutant forms of genes encoding minK protein and KVLQT1 protein involved
 PT in cardiac potassium channel formation useful for screening drugs, for
 PT preventing and treating cardiac arrhythmia -
 XX
 PS Disclosure; Fig 3; 167pp; English.
 XX
 CC The invention relates to KVLQT1 and KCNE1 genes, associated with long
 CC QT (LQT) syndrome. It provides a minK protein comprising a mutation which
 CC substitutes the wild type amino acids with Leu, Asp, Leu, His, Trp and
 CC Ala or Thr at residues 74, 76, 28, 32, 98 and 127 respectively. Screening
 CC KVLQT1 and KCNE1 is useful for identifying mutations for diagnosing and
 CC treating LQT. The ability to predict LQT enables physicians to prevent
 CC the diseases with medical therapy such as beta blocking agents and opts
 CC for better treatments. The present sequence represents the human
 CC KVLQT1 protein fragment.
 XX
 SQ Sequence 137 AA;
 Query Match 18.4%; Score 69; DB 21; Length 137;
 Best Local Similarity 100.0%; Pred. No. 1.8e-59;
 Matches 69; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 185 GOVFATSAIRGIRFLOILRMLHVDROGQWRLGSGVVFHROELITTLTYIGFLGIFSSY 244
 DB 13 gqvfatSaIRGIRfLQILrmlhvdrggqtwrllsgsvvfhrqelittlyigflglifssy 62
 QY 245 FVYLAEKDA 253
 DB 63 fvyLaekda 71
 RESULT 5
 AAY57377
 ID AAY57377 standard; Protein; 570 AA.
 XX
 AC AAY57377;
 XX
 DT 19-JUN-2000 (first entry)
 XX
 DE Human KVLQT1 protein fragment.
 XX
 KW KVLQT1; KCNE1; long QT syndrome; LQT syndrome; minK protein;
 KW antiarrhythmic; gene therapy; human.
 XX

PS Claim 24; Page 72-73; 105pp; English.

XX This polypeptide comprises the xenopus homologue of human KVLQT1

CC (see AAW30038), a protein associated with long QT syndrome (LQT). A

CC cDNA clone encoding xenopus KVLQT1 was isolated from an oocyte cDNA

CC library by homology to human KVLQT1 cDNA (see AAT90730). Human

CC KVLQT1 coassembles with human minK to form a cardiac IKs potassium

CC channel. Coexpression of these proteins in a cell can be used to

CC screen for drugs useful in treating or preventing LQT.

XX Sequence 376 AA;

Query Match 100.0%; Score 376; DB 18; Length 376;

Best Local Similarity 100.0%; Pred. No. 0;

Matches 376; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MNENAINSLYEALPLPDGSSNGQRQEDRQANSFELKRETLVATDPPRPTINLDPVRSIY 60

Db 1 mnenainslyealplpdgssngqrqedrqansfelkretlvdpprptinldprvsiy 60

QY 61 SGRPLFSRTNIQGRVYNFLRPTGKCFVYHFTVFLVLICLIFSLSVSTIQYNNLATE 120

Db 61 sgrplfstrtniqgrvynflerptgkcfvyhftvflvlliclifsylvstiqgynnlata 120

QY 121 TLFWMEIVLVVFFGAAYVVRWSAGCRSKYGVWGRRLSFARKPIVSIDLIWVASVIVLC 180

Db 121 tlfwmeivlvvffgaeyvvrwlsagcrskylvwgrlrfarkpislvidliwvasvivlc 180

QY 181 VGSNGQVFATSAIRGIRFLQILRMLHVDROGTTWRLLSGVVFIHQELITTLTIGFLGLI 240

Db 181 vgsngqvfatalsairgrfqlrlmlhvdrgttwrlsgvvfihqelittltigflgli 240

QY 241 FSSYFVYLAEKDAIDSSEYQFGSYADALWVGWVTVTTIGYGDVPOTWIGKTIASCFSV 300

Db 241 fssyfvyllaekdaidsseeyqfgyadallwvgwvttvttigygdkvpotwigktiascfsv 300

QY 301 FAISFFALPAGILSGFALKVQKQKQKHFNRQIPAAASLIQTAWRCYAAENPDATWKI 360

Db 301 faisffalpagilsgfalkvqkqkqkhnrfqpaaasliqtawrcyaaenpdatswki 360

QY 361 YIRKQSRNHHIMSPSP 376

Db 361 yirkqsrnhhimspsp 376

RESULT 2

AAAY57376

ID AAY57376 standard; Protein; 376 AA.

XX AAY57376;

XX 19-JUN-2000 (first entry)

DE xenopus KVLQT1 partial protein fragment.

XX KVLQT1; KCNE1; long QT syndrome; LQT syndrome; minK protein;

KW antiarrhythmic; gene therapy; human; frog.

XX xenopus laevis.

XX WO200006600-A1.

XX 10-FEB-2000.

XX 06-OCT-1998; 98WO-US17838.

XX 29-JUL-1998; 98US-0094477.

PR 17-AUG-1998; 98US-0135020.

XX (UTAH) UNIV UTAH RES FOUND.

XX (Sanguinetti MC, Splawski I;

PI

XX WPI; 2000-195262/17.

XX Mutant forms of genes encoding minK protein and KVLQT1 protein involved

PT in cardiac potassium channel formation useful for screening drugs, for

PT preventing and treating cardiac arrhythmia

XX Disclosure; Fig 10; 167pp; English.

XX The invention relates to KVLQT1 and KCNE1 genes, associated with long

CC QT (LQT) syndrome. It provides a minK protein comprising a mutation which

CC substitutes the wild type amino acids with Leu, Asp, Leu, His, Trp and

CC Ala or Thr at residues 74,76,28,32,98 and 127 respectively. Screening

CC KVLQT1 and KCNE1 is useful for identifying mutations for diagnosing and

CC treating LQT. The ability to predict LQT enables physicians to prevent

CC the diseases with medical therapy such as beta blocking agents and opts

CC for better treatments. The present sequence represents a Xenopus

CC KVLQT1 partial protein fragment.

XX Sequence 376 AA;

Query Match 100.0%; Score 376; DB 21; Length 376;

Best Local Similarity 100.0%; Pred. No. 0;

Matches 376; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MNENAINSLYEALPLPDGSSNGQRQEDRQANSFELKRETLVATDPPRPTINLDPVRSIY 60

Db 1 mnenainslyealplpdgssngqrqedrqansfelkretlvdpprptinldprvsiy 60

QY 61 SGRPLFSRTNIQGRVYNFLRPTGKCFVYHFTVFLVLICLIFSLSVSTIQYNNLATE 120

Db 61 sgrplfstrtniqgrvynflerptgkcfvyhftvflvlliclifsylvstiqgynnlata 120

QY 121 TLFWMEIVLVVFFGAAYVVRWSAGCRSKYGVWGRRLSFARKPIVSIDLIWVASVIVLC 180

Db 121 tlfwmeivlvvffgaeyvvrwlsagcrskylvwgrlrfarkpislvidliwvasvivlc 180

QY 181 VGSNGQVFATSAIRGIRFLQILRMLHVDROGTTWRLLSGVVFIHQELITTLTIGFLGLI 240

Db 181 vgsngqvfatalsairgrfqlrlmlhvdrgttwrlsgvvfihqelittltigflgli 240

QY 241 FSSYFVYLAEKDAIDSSEYQFGSYADALWVGWVTVTTIGYGDVPOTWIGKTIASCFSV 300

Db 241 fssyfvyllaekdaidsseeyqfgyadallwvgwvttvttigygdkvpotwigktiascfsv 300

QY 301 FAISFFALPAGILSGFALKVQKQKQKHFNRQIPAAASLIQTAWRCYAAENPDATWKI 360

Db 301 faisffalpagilsgfalkvqkqkqkhnrfqpaaasliqtawrcyaaenpdatswki 360

QY 361 YIRKQSRNHHIMSPSP 376

Db 361 yirkqsrnhhimspsp 376

RESULT 3

AAAY80567

ID AAY80567 standard; Protein; 376 AA.

XX AAY80567;

XX 06-JUN-2000 (first entry)

DE partial Xenopus KVLQT1 protein.

XX KVLQT1; mutation; human; cardiac I(ks) potassium channel; KCNE1; ss;

KW cardiac arrhythmia; electrocardiogram; Long QT syndrome; gene therapy.

XX xenopus laevis.

XX WO200006199-A1.

XX 10-FEB-2000.

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: November 3, 2001, 10:57:35 ; Search time 33.71 Seconds
(without alignments)
676.198 Million cell updates/sec

Title: US-09-135-010A-113
Perfect score: 376
Sequence: 1 MNEAINSLYRAIPDPDGS.....TWKIYKQSRNHHIMSPSP 376

Scoring table: OLIGO
Gapop 60.0 , Gapext 60.0

Searched: 412676 seqs, 60623988 residues

Word size : 4

Total number of hits satisfying chosen parameters: 121002

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : A_Geneseq_0601.*
1: /SIDS1/gcgdata/geneseq/geneseq/AA1980.DAT.*
2: /SIDS1/gcgdata/geneseq/geneseq/AA1981.DAT.*
3: /SIDS1/gcgdata/geneseq/geneseq/AA1982.DAT.*
4: /SIDS1/gcgdata/geneseq/geneseq/AA1983.DAT.*
5: /SIDS1/gcgdata/geneseq/geneseq/AA1984.DAT.*
6: /SIDS1/gcgdata/geneseq/geneseq/AA1985.DAT.*
7: /SIDS1/gcgdata/geneseq/geneseq/AA1986.DAT.*
8: /SIDS1/gcgdata/geneseq/geneseq/AA1987.DAT.*
9: /SIDS1/gcgdata/geneseq/geneseq/AA1988.DAT.*
10: /SIDS1/gcgdata/geneseq/geneseq/AA1989.DAT.*
11: /SIDS1/gcgdata/geneseq/geneseq/AA1990.DAT.*
12: /SIDS1/gcgdata/geneseq/geneseq/AA1991.DAT.*
13: /SIDS1/gcgdata/geneseq/geneseq/AA1992.DAT.*
14: /SIDS1/gcgdata/geneseq/geneseq/AA1993.DAT.*
15: /SIDS1/gcgdata/geneseq/geneseq/AA1994.DAT.*
16: /SIDS1/gcgdata/geneseq/geneseq/AA1995.DAT.*
17: /SIDS1/gcgdata/geneseq/geneseq/AA1996.DAT.*
18: /SIDS1/gcgdata/geneseq/geneseq/AA1997.DAT.*
19: /SIDS1/gcgdata/geneseq/geneseq/AA1998.DAT.*
20: /SIDS1/gcgdata/geneseq/geneseq/AA1999.DAT.*
21: /SIDS1/gcgdata/geneseq/geneseq/AA2000.DAT.*
22: /SIDS1/gcgdata/geneseq/geneseq/AA2001.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	376	100.0	376	18 AAW30036	Xenopus KVLQT1.. X
2	376	100.0	376	21 AAY57376	Xenopus KVLQT1 par
3	376	100.0	376	21 AAY80567	partial Xenopus KV
4	69	18.4	137	21 AAY57372	Human KVLQT1 prote
5	69	18.4	570	21 AAY57377	Human KVLQT1 prote
6	69	18.4	581	18 AAW33355	Human KVLQT1 assoc
7	69	18.4	581	18 AAW30038	Human KVLQT1 assoc
8	69	18.4	581	22 AAB49499	Mutant human KVLQT
9	69	18.4	676	21 AAY57368	Human KVLQT1 prote
10	69	18.4	676	21 AAY80562	Human long QT synd
11	69	18.4	676	22 AAB49494	Human KVLQT1.. Hom

12	69	18.4	677	20	AAW08343	Human KCNQ1 protei
13	21	5.6	21	21	AAB11385	Potassium channel
14	21	5.6	283	22	AAB49495	Mutant human KVLQT
15	21	5.6	807	20	AAW08342	Human nKQT1 protei
16	20	5.3	695	21	AAB01476	KCNQ4 Potassium ch
17	20	5.3	846	21	AAB24241	Human KCNQ5 (KCN6g
18	20	5.3	897	22	AAB47046	Human KCNQ5 potass
19	19	5.1	245	20	AAW01531	Amino acid sequenc
20	19	5.1	393	18	AAW14282	Human K+ channel p
21	19	5.1	722	20	AAW01530	Amino acid sequenc
22	19	5.1	757	20	AAW08345	Mouse partial KCNQ
23	19	5.1	854	20	AAW23215	Human brain-derive
24	19	5.1	871	20	AAW01529	Amino acid sequenc
25	19	5.1	872	20	AAW08341	Human KCNQ2 protei
26	19	5.1	930	20	AAW08347	Human mutant KCNQ2
27	17	4.5	61	21	AAW57371	Human KVLQT1 prote
28	15	4.0	15	20	AAW34136	Variant human pota
29	13	3.5	854	20	AAW01534	Amino acid sequenc
30	13	3.5	870	20	AAW08346	Mouse KCNQ3 protei
31	13	3.5	872	20	AAW08344	Human KCNQ3 protei
32	10	2.7	26	21	AAW57370	Human KVLQT1 prote
33	8	2.1	1600	21	AAW51095	Arabidopsis thalia
34	8	2.1	1608	21	AAW51094	Arabidopsis thalia
35	8	2.1	1625	21	AAW51093	Arabidopsis thalia
36	7	1.9	14	15	AAW52597	Hepatitis E virus
37	7	1.9	67	20	AAW11640	Human 5' EST seque
38	7	1.9	74	20	AAW35977	Extended human sec
39	7	1.9	98	21	AAW16753	Bacteriophage Dp-1
40	7	1.9	119	19	AAW69227	NADH dehydrogenase
41	7	1.9	143	21	AAW54054	Angiotensin-bindin
42	7	1.9	261	20	AAW06553	Phage lambda red b
43	7	1.9	275	15	AAW60476	Serine protease of
44	7	1.9	327	17	AAW96093	Hepatitis E virus
45	7	1.9	327	18	AAW35820	Hepatitis E virus

ALIGNMENTS

RESULT 1
AAW30036
ID AAW30036 standard; protein; 376 AA.
XX
AC AAW30036;
XX
DT 12-FEB-1998 (first entry)
DT
XX
DE Xenopus KVLQT1.
XX
KW KVLQT1; long QT syndrome; arrhythmia; mink; potassium channel;
KW diagnosis; therapy.
XX
OS Xenopus sp.
XX
PN WO9723598-A2.
XX
PD 03-JUL-1997.
XX
PF 20-DEC-1996; 96WO-US19756.
XX
PR 29-OCT-1996; 96US-0739383.
PR 22-DEC-1995; 95US-0019014.
XX
PA (UTAH) UNIV UTAH RES FOUND.
XX
PI Curran ME, Keating MT, Sanguinetti MC;
XX
DR WPI; 1997-402190/37.
XX
PT Human mink and Xenopus KVLQT1 coding sequences - used for assays for
PT identifying drugs which can be used for preventing or treating long
PT QT syndrome
XX

Best Local Similarity 100.0%; Pred. No. 22;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 196 IRLQL 202
|||||
Db 13 IRLQL 19

RESULT 15

TBPA_HAEIN STANDARD; PRT; 332 AA.
AC P44984;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 01-OCT-2000 (Rel. 40, Last annotation update)
DE THIAMINE-BINDING PERIPLASMIC PROTEIN PRECURSOR.
GN TBPA OR H11019.
OS Haemophilus influenzae.
OC Bacteria; Proteobacteria; gamma subdivision; Pasteurellaceae;
OC Haemophilus.
OX NCBI_TaxID=727;
RN [1]

RP SEQUENCE FROM N.A.
RC STRAIN=RD / KW20 / ATCC 51907;
RX MEDLINE=95350630; PubMed=7542800;
RA Frelschiemann R.D., Adams M.D., White O., Clayton R.A., Kirkness E.F.,
RA Kerlavage A.R., Bult C.J., Tomb J.-F., Dougherty B.A., Merrick J.M.,
RA McKenney K., Sutton G., Fitzhugh W., Fields C.A., Gocayne J.D.,
RA Scott J.D., Shirley R., Liu L.-I., Glodek A., Kelley J.M.,
RA Weidman J.F., Phillips C.A., Spriggs T., Hedblom E., Cotton M.D.,
RA Utterback T.R., Hanna M.C., Nguyen D.T., Saudek D.M., Brandon R.C.,
RA Fine L.D., Fritchman J.L., Fuhrmann J.L., Geoghagen N.S.M.,
RA Gnehm C.L., McDonald L.A., Small K.V., Fraser C.M., Smith H.O.,
RA Venter J.C.;
RT "Whole-genome random sequencing and assembly of Haemophilus
RT influenzae Rd";
RL Science 269:496-512(1995).
CC -!- FUNCTION: PART OF THE BINDING-PROTEIN-DEPENDENT TRANSPORT SYSTEM
CC TBPA-THIPIQ FOR THIAMINE AND TPP (BY SIMILARITY).
CC -!- SUBCELLULAR LOCATION: PERIPLASMIC.
CC -!- SIMILARITY: BELONGS TO THE BACTERIAL EXTRACELLULAR SOLUTE-BINDING
CC PROTEIN FAMILY 1.

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DR EMBL: U32782; AAC22678.1; -
DR TIGR: H11019; -
DR InterPro: IPR000567; -
DR PROSITE: PS01037; SBP_BACTERIAL_1; 1.
KW Transport; periplasmic; Signal.
FT SIGNAL 1 20 POTENTIAL.
FT CHAIN 21 332 THIAMINE-BINDING PERIPLASMIC PROTEIN.
SQ SEQUENCE 332 AA: 37272 MW: 90A27B35D0F9C741 CRC64;

Query Match 1.9%; Score 7; DB 1; Length 332;
Best Local Similarity 100.0%; Pred. No. 27;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 318 ALKVOOK 324
|||||
Db 299 ALKVOOK 305

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DR EMBL: J02058; -; NOT_ANNOTATED_CDS.
DR PIR: A04168; QQOMC2.
DR InterPro: IPR000263; -.
DR InterPro: IPR001530; -.
DR InterPro: IPR003001; -.
DR Pfam: PF01489; Gemini_BR1.1.
DR PRINTS: PR00223; GEMCONTARBR1.
DR PRINTS: PR00225; GEMCONTBR1.
SQ SEQUENCE 256 AA; 29305 MW; 070A364569507634 CRC64;

Query Match 1.9%; Score 7; DB 1; Length 256;
Best Local Similarity 100.0%; Pred. No. 22;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 362 IRKOSRN 368
DB 4 IRKOSRN 10

RESULT 13
-VBET_LAMB
ID VBET_LAMB STANDARD; PRT; 261 AA.
AC P03698;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE RECOMBINATION PROTEIN BET
GN BET OR BETA OR RED-BETA OR REDB.
OS Bacteriophage lambda.
OC Viruses; dsDNA viruses, no RNA stage; Tailed phages; Siphoviridae;
OC Lambda phage group.
OX NCBI_TaxID=10710;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=83189071; PubMed=6221115;
RA Sanger F., Coulson A.R., Hong G.F., Hill D.F., Petersen G.B.;
RT "Nucleotide sequence of bacteriophage lambda DNA.";
RL J. Mol. Biol. 162:729-773(1982).
RN [2]
RP SEQUENCE OF 1-103 FROM N.A.
RX MEDLINE=82059489; PubMed=6458018;
RA Ineichen K., Shepherd J.C.W., Bickle T.A.;
RT "The DNA sequence of the phage lambda genome between PL and the gene
RT bet.";
RL Nucleic Acids Res. 9:4639-4653(1981).
CC -!- FUNCTION: GENE BET PROTEIN FUNCTIONS IN GENERAL RECOMBINATION AND
CC IN THE LATE, ROLLING-CIRCLE MODE OF LAMBDA DNA REPLICATION.
CC HAS A FUNCTION SIMILAR TO THAT OF E.COLI RECT. IT IS A SINGLE-
CC STRANDED DNA BINDING PROTEIN THAT CAN PROMOTE RENATURATION OF DNA.

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DR EMBL: J02459; AAA96570.1; -.
DR EMBL: V00638; CAA23976.1; -.
DR PIR: A04320; QBBPL.
KW DNA recombination; DNA-binding.
SQ SEQUENCE 261 AA; 29688 MW; 99583014F97330A6 CRC64;

CC Query Match 1.9%; Score 7; DB 1; Length 261;
CC Best Local Similarity 100.0%; Pred. No. 22;
CC Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 226 QELITTL 232
DB 24 QELITTL 30

RESULT 14
PLSC_MYCGE
ID PLSC_MYCGE STANDARD; PRT; 268 AA.
AC Q49402; Q49287;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE PROBABLE 1-ACYL-SN-GLYCEROL-3-PHOSPHATE ACYLTRANSFERASE (EC 2.3.1.51)
DE (1-AGP ACYLTRANSFERASE) (1-AGPAT) (LYSOPHOSPHATIDIC ACID
DE ACYLTRANSFERASE) (LPAAT).
GN PLSC OR MG212.
OS Mycoplasma genitalium.
OC Bacteria; Firmicutes; Bacillus/Clostridium group; Mollicutes;
OC Mycoplasmataceae; Mycoplasma.
OX NCBI_TaxID=2097;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 33530 / G-37;
RX MEDLINE=96026346; PubMed=7569993;
RA Fraser C.M., Gocayne J.D., White O., Adams M.D., Clayton R.A.,
RA Fleischmann R.D., Bult C.J., Kerlavage A.R., Sutton G., Kelley J.M.,
RA Fritchman J.L., Weidman J.F., Small K.V., Sandusky M., Fuhrmann J.L.,
RA Nguyen D.T., Utterback T.R., Saudek D.M., Phillips C.A., Merrick J.M.,
RA Tomb J.-F., Dougherty B.A., Bott K.F., Hu P.-C., Lucier T.S.,
RA Peterson S.N., Smith H.O., Hutchison C.A. III, Venter J.C.;
RT "The minimal gene complement of Mycoplasma genitalium.";
RL Science 270:397-403(1995).
RN [2]
RP SEQUENCE OF 1-105 FROM N.A.
RC STRAIN=ATCC 33530 / G-37;
RX MEDLINE=94075230; PubMed=8253680;
RA Peterson S.N., Hu P.-C., Bott K.F., Hutchison C.A. III;
RT "A survey of the Mycoplasma genitalium genome by using random
RT sequencing.";
RL J. Bacteriol. 175:7918-7930(1993).
CC -!- FUNCTION: CONVERTS LYSOPHOSPHATIDIC ACID (LPA) INTO PHOSPHATIDIC
CC -!- ACID BY INCORPORATING ACYL MOIETY AT THE 2 POSITION.
CC -!- CATALYTIC ACTIVITY: ACYL-COA + 1-ACYL-SN-GLYCEROL 3-PHOSPHATE -
CC COA + 1,2-DIACYL-SN-GLYCEROL 3-PHOSPHATE.
CC -!- PATHWAY: SECOND STEP IN DE NOVO PHOSPHOLIPID BIOSYNTHESIS.
CC -!- SIMILARITY: BELONGS TO THE 1-ACYL-SN-GLYCEROL-3-PHOSPHATE
CC ACYLTRANSFERASE FAMILY.

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DR EMBL: U39701; AAC71431.1; -.
DR EMBL: U02160; AAD12442.1; -.
DR TIGR: MG212; -.
DR InterPro: IPR002123; -.
KW Phospholipid biosynthesis; Transferrase; Acyltransferase.
FT CONFLICT 2 MISSING (IN REF. 2).
SQ SEQUENCE 268 AA; 30469 MW; A88B07D2BC4C6A4A CRC64;

Query Match 1.9%; Score 7; DB 1; Length 268;

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FT DOMAIN 122 139 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 140 160 POTENTIAL.
FT DOMAIN 161 177 LUMENAL (POTENTIAL).
FT BINDING 147 147 DICYCLOHEXYLCARODIIMIDE (POTENTIAL).
SQ SEQUENCE 177 AA; 18131 MW; 32521191B721FB52 CRC64;

Query Match 1.9%; Score 7; DB 1; Length 177;
Best Local Similarity 100.0%; Pred. No. 16;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 100 LICLIIFS 106
Db 140 LICLIIFS 146

RESULT 10
ID Y212.METJA STANDARD; PRT; 241 AA.
AC Q60274;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE HYPOTHETICAL PROTEIN MJECL12.
GN MJECL12.
OS Methanococcus jannaschii.
OC Archaea; Euryarchaeota; Methanococcales; Methanococcaceae;
OC Methanococcus.
OX NCBI_TaxID=2190;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=JAL-1 / DSM 2661 / ATCC 43067;
RX MEDLINE=96337999; PubMed=8688087;
RA Bult C.J., White O., Olsen G.J., Zhou L., Fleischmann R.D.,
Sutton G.G., Blake J.A., Fitzgerald L.M., Clayton R.A., Gocayne J.D.,
Kerlavage A.R., Dougherty B.A., Tomb J.F., Adams M.D., Reich C.I.,
Overbeek R., Kirkness E.F., Weinstock K.G., Merrick J.M., Glodek A.,
Scott J.L., Geoghegan N.S.M., Weidman J.F., Fuhrmann J.L., Nguyen D.,
Utterback T.R., Kelley J.M., Peterson J.D., Sadow P.W., Hanna M.C.,
Cotton M.D., Roberts K.M., Hurst M.A., Kaine B.P., Borodovsky M.,
Klenk H.-P., Fraser C.M., Smith H.O., Woese C.R., Venter J.C.;
"Complete genome sequence of the methanogenic archaeon, Methanococcus
jannaschii."
RL Science 273:1058-1073(1996).
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CC -----
DR EMBL; L77118; AAC37085.1; -
DR TIGR; MJECL12; -
KW Hypothetical protein.
SQ SEQUENCE 241 AA; 27268 MW; C5D3C7742A35A097 CRC64;

Query Match 1.9%; Score 7; DB 1; Length 241;
Best Local Similarity 100.0%; Pred. No. 21;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 33 SFELKRE 39
Db 148 SFELKRE 154

RESULT 11
ATP6_BACSU
ID ATP6_BACSU STANDARD; PRT; 244 AA.
AC P37813;
DT 01-OCT-1994 (Rel. 30, Created)

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DT 01-OCT-1994 (Rel. 30, Last sequence update)
DT 01-NOV-1995 (Rel. 32, Last annotation update)
DE ATP SYNTHASE A CHAIN (EC 3.6.1.34) (PROTEIN 6).
GN ATPB.
OS Bacillus subtilis.
OC Bacteria; Firmicutes; Bacillus/Clostridium group;
OC Bacillus/Staphylococcus group; Bacillus.
OX NCBI_TaxID=1423;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=168;
RX MEDLINE=95050246; PubMed=7961438;
RA Santana M., Ionescu M.S., Vertes A., Longin R., Kunst F., Danchin A.,
Glaser P.;
RT "Bacillus subtilis F01 ATPase: DNA sequence of the atp operon and
characterization of atp mutants."
RL J. Bacteriol. 176:6802-6811(1994).
CC -----
CC -1- FUNCTION: NOT COMPONENT OF THE PROTON CHANNEL: IT MAY PLAY A
DIRECT ROLE IN THE TRANSLOCATION OF PROTONS ACROSS THE MEMBRANE.
CC -1- SUBUNIT: F-TYPE ATPASES HAVE 2 COMPONENTS, CF(1) - THE CATALYTIC
CORE - AND CF(0) - THE MEMBRANE PROTON CHANNEL. CF(1) HAS FIVE
SUBUNITS: ALPHA(3), BETA(3), GAMMA(1), DELTA(1), EPSILON(1). CF(0)
HAS THREE MAIN SUBUNITS: A, B AND C.
CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN. CONTAINS 8
POTENTIAL TRANSMEMBRANE DOMAINS
CC -1- SIMILARITY: BELONGS TO THE ATPASE A CHAIN FAMILY.
CC -----
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or send an email to license@isb-sib.ch).
CC -----
DR EMBL; Z28592; CAA82254.1; -
DR EMBL; Z99122; CAB15704.1; -
DR PIR; S39250; S39250.
DR Subtilist; BG10815; atpB.
DR InterPro; IPR000568; -
DR Pfam; PF00119; ATP-synt_A; 1.
DR PROSITE; PS00449; ATPASE_A; 1.
KW Hydrogen ion transport; CF(0); Transmembrane.
SQ SEQUENCE 244 AA; 27054 MW; E26172BA9F1AA248 CRC64;

Query Match 1.9%; Score 7; DB 1; Length 244;
Best Local Similarity 100.0%; Pred. No. 21;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 173 VASVIVL 179
Db 25 VASVIVL 31

RESULT 12
ID VBR1_CLVK STANDARD; PRT; 256 AA.
AC Q93565;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 01-JUN-1994 (Rel. 29, Last annotation update)
DE BRL PROTEIN (29.4 KDA PROTEIN).
GN BVL.
OS Cassava latent virus (strain West Kenyan 844).
OC Viruses; ssDNA viruses; Geminiviridae; Begomovirus.
OX NCBI_TaxID=10818;
RN [1]
RP SEQUENCE FROM N.A.
RA Stanley J., Gay M.R.;
RT "Nucleotide sequence of cassava latent virus DNA."
RL Nucleic Acids Res. 11:260-262(1983).
CC -----
CC -1- SIMILARITY: BELONGS TO GEMINIVIRUSES BRL PROTEIN FAMILY.

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DR PROSITE: PS00598; CHROMO.1; 1.
DR PROSITE: PS00113; CHROMO.2; 2.
KW Chromatin regulator; Nuclear protein; Transcription regulation;
KW Repressor; Phosphorylation.
FT DOMAIN 20 78 CHROMO.
FT DOMAIN 111 169 CHROMO SHADOW DOMAIN.
SQ SEQUENCE 173 AA; 19720 MW; EB9D2F554F58C897 CRC64;

Query Match 1.9%; Score 7; DB 1; Length 173;
Best Local Similarity 100.0%; Pred. No. 16;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 253 AIDSSGE 259
|||||||
DB 119 AIDSSGE 125

RESULT 8
VATL_ENTDI
ID VATL_ENTDI STANDARD; PRT; 176 AA.
AC Q24808;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE VACUOLAR ATP SYNTHASE 16 KDA PROTEOLIPID SUBUNIT (EC 3.6.1.34) (V-
ATPASE 16 KDA PROTEOLIPID SUBUNIT).
GN VMA3.
OS Entamoeba dispar.
OC Eukaryota; Entamoebidae; Entamoeba.
OX NCBI_TaxID=46681;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=NON-PATHOGENIC;
RX MEDLINE=94314485; PubMed=8039932;
RA Descoeurs S., Yu Y., Samuelson J.;
RT "Cloning of Entamoeba genes encoding proteolipids of putative
vaccuolar proton-translocating ATPases.";
RL Infect. Immun. 62:3572-3575(1994).
CC -!- FUNCTION: PROTON-CONDUCTING PORE FORMING SUBUNIT OF THE MEMBRANE
INTEGRAL V0 COMPLEX OF VACUOLAR ATPASE. V-ATPASE IS RESPONSIBLE
FOR ACIDIFYING A VARIETY OF INTRACELLULAR COMPARTMENTS IN
EUKARYOTIC CELLS.
CC -!- SUBUNIT: V-ATPASE IS AN HETEROMULTIMERIC ENZYME COMPOSED OF A
PERIPHERAL CATALYTIC V1 COMPLEX (MAIN COMPONENTS: SUBUNITS A, B,
C, D, E, AND F) ATTACHED TO AN INTEGRAL MEMBRANE V0 PROTON PORE
COMPLEX (MAIN COMPONENT: THE PROTEOLIPID PROTEIN; WHICH IS PRESENT
AS A HEXAMER THAT FORMS THE PROTON-CONDUCTING PORE).
CC -!- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN. VACUOLAR.
CC -!- MISCELLANEOUS: THIS SUBUNIT BINDS DICYCLOHEXYLCARBODIIMIDE (DCDD)
WHICH INHIBITS THE ATPASE (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE V-ATPASE PROTEOLIPID SUBUNIT FAMILY.

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EMBL; U01055; AAA21448.1; -
InterPro; IPR000245; -
DR InterPro; IPR002379; -
DR Pfam; PF00137; ATP-synt_C; 2.
DR PRINTS; PR00122; VACATPASE.
KW Hydrolase; Hydrogen ion transport; Transmembrane.
FT DOMAIN 1 17 LUMENAL (POTENTIAL).
FT TRANSMEM 18 38 POTENTIAL.
FT DOMAIN 39 62 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 63 83 POTENTIAL.
FT DOMAIN 84 98 LUMENAL (POTENTIAL).
FT TRANSMEM 99 119 POTENTIAL.

FT DOMAIN 120 136 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 137 157 POTENTIAL.
FT DOMAIN 158 176 LUMENAL (POTENTIAL).
FT BINDING 145 145 DICYCLOHEXYLCARBODIIMIDE (POTENTIAL).
FT SITE 145 145 ESSENTIAL FOR ENZYME AND TRANSPORT
ACTIVITY (BY SIMILARITY).
SQ SEQUENCE 176 AA; 18103 MW; 50132CC98FD0B850 CRC64;

Query Match 1.9%; Score 7; DB 1; Length 176;
Best Local Similarity 100.0%; Pred. No. 16;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 100 LICLIIPS 106
|||||||
DB 138 LICLIIPS 144

RESULT 9
VATL_ENTHI
ID VATL_ENTHI STANDARD; PRT; 177 AA.
AC Q24810;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE VACUOLAR ATP SYNTHASE 16 KDA PROTEOLIPID SUBUNIT (EC 3.6.1.34) (V-
ATPASE 16 KDA PROTEOLIPID SUBUNIT).
GN VMA3.
OS Entamoeba histolytica.
OC Eukaryota; Entamoebidae; Entamoeba.
OX NCBI_TaxID=5759;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=HM-1; IMSS;
RX MEDLINE=94314485; PubMed=8039932;
RA Descoeurs S., Yu Y., Samuelson J.;
RT "Cloning of Entamoeba genes encoding proteolipids of putative
vaccuolar proton-translocating ATPases.";
RL Infect. Immun. 62:3572-3575(1994).
CC -!- FUNCTION: PROTON-CONDUCTING PORE FORMING SUBUNIT OF THE MEMBRANE
INTEGRAL V0 COMPLEX OF VACUOLAR ATPASE. V-ATPASE IS RESPONSIBLE
FOR ACIDIFYING A VARIETY OF INTRACELLULAR COMPARTMENTS IN
EUKARYOTIC CELLS.
CC -!- SUBUNIT: V-ATPASE IS AN HETEROMULTIMERIC ENZYME COMPOSED OF A
PERIPHERAL CATALYTIC V1 COMPLEX (MAIN COMPONENTS: SUBUNITS A, B,
C, D, E, AND F) ATTACHED TO AN INTEGRAL MEMBRANE V0 PROTON PORE
COMPLEX (MAIN COMPONENT: THE PROTEOLIPID PROTEIN; WHICH IS PRESENT
AS A HEXAMER THAT FORMS THE PROTON-CONDUCTING PORE).
CC -!- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN. VACUOLAR.
CC -!- MISCELLANEOUS: THIS SUBUNIT BINDS DICYCLOHEXYLCARBODIIMIDE (DCDD)
WHICH INHIBITS THE ATPASE (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE V-ATPASE PROTEOLIPID SUBUNIT FAMILY.

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EMBL; U01057; AAA21450.1; -
InterPro; IPR000245; -
DR InterPro; IPR002379; -
DR Pfam; PF00137; ATP-synt_C; 2.
DR PRINTS; PR00122; VACATPASE.
KW Hydrolase; Hydrogen ion transport; ATP synthesis; Transmembrane.
FT DOMAIN 1 19 LUMENAL (POTENTIAL).
FT TRANSMEM 20 40 POTENTIAL.
FT DOMAIN 41 64 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 65 85 POTENTIAL.
FT DOMAIN 86 100 LUMENAL (POTENTIAL).
FT TRANSMEM 101 121 POTENTIAL.

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SQ SEQUENCE 119 AA: 14187 MW: 9A47DEB33DC9244D CRC64;

Query Match 1.9%; Score 7; DB 1; Length 119;
Best Local Similarity 100.0%; Pred. No. 11;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 232 LVIGFLG 238
      |||||
Db 31 LVIGFLG 37

RESULT 7
CBX3_HUMAN
ID ID CBX3_HUMAN STANDARD; PRT; 173 AA.
Q13185; Q99409;
DT 01-NOV-1997 (Rel. 35, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 01-OCT-2000 (Rel. 40, Last annotation update)
DE CHROMOX PROTEIN HOMOLOG 3 (HETEROCHROMATIN PROTEIN 1 HOMOLOG GAMMA)
DE (HPI GAMMA) (MODIFIER 2 PROTEIN).
GN CBX3.
OS Homo sapiens (Human)
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
NCBI_TaxID=9606;
RN [1].
RP SEQUENCE FROM N.A.
RX MEDLINE=96278941; PubMed=8663349;
RA Ye Q., Workman H.J.;
RT "Interaction between an integral protein of the nuclear envelope
RT inner membrane and human chromodomain proteins homologous to
RT Drosophila HPI.";
RT J. Biol. Chem. 271:14653-14656(1996).
RN [2].
RP REVISIONS.
RA Ye Q., Workman H.J.;
RRL Submitted (JAN-1997) to the EMBL/GenBank/DBJ databases.
RN [3].
RP SEQUENCE FROM N.A.
RX MEDLINE=20130009; PubMed=10664448;
RA Koike N., Maita H., Taira T., Ariga H., Iguchi-Ariga S.M.M.;
RT "(d)entification of heterochromatin protein 1 (HPI) as a
RT phosphorylation target by pim-1 kinase and the effect of
RT phosphorylation on the transcriptional repression function of HPI.";
RT FEBS Lett. 467:17-21(2000).
CC !- FUNCTION: COMPONENT OF HETEROCHROMATIN. MAY INTERACT WITH LAMIN B
CC RECEPTOR (LBR). THIS INTERACTION CAN CONTRIBUTE TO THE ASSOCIATION
CC OF THE HETEROCHROMATIN WITH THE INNER NUCLEAR MEMBRANE.
CC !- SUBCELLULAR LOCATION: NUCLEAR (POTENTIAL).
CC !- PTM: PHOSPHORYLATION OF HPI AND LBR MAY BE RESPONSIBLE FOR SOME OF
CC THE ALTERATIONS IN CHROMATIN ORGANIZATION AND NUCLEAR STRUCTURE
CC WHICH OCCUR AT VARIOUS TIMES DURING THE CELL CYCLE. PHOSPHORYLATED
CC BY PIM-1.
CC !- SIMILARITY: CONTAINS 1 'CHROMO' DOMAIN AND 1 'CHROMO SHADOW'
CC DOMAIN.
CC
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CC
CC EMBL; U26312; AAB48101.1; -.
CC EMBL; AB030905; BAA83340.1; -.
CC MIM; 604477; -.
CC HSSP; Z23197; IAP0.
CC InterPro; IPR000953; -.
CC Pfam; PF01393; Chromo_shadow; 1.
CC Pfam; PF00385; chromo; 1.
CC PRINTS; PR00504; CHROMODOMAIN.

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DR EMBL; AF105212; AAD14681.1; JOINED.
DR EMBL; AF105213; AAD14681.1; JOINED.
DR EMBL; AF105214; AAD14681.1; JOINED.
DR EMBL; AF105215; AAD14681.1; JOINED.
DR MIM; 603537; -.
DR MIM; 600101; -.
DR InterPro; IPR000636; -.
DR InterPro; IPR003091; -.
DR Pfam; PF00520; Ion_trans; 1.
DR PRINTS; PR00169; KCHANNEL. 6
KW Ionic channel; Transmembrane; Ion transport; Voltage-gated channel;
KW Multigene family; Disease mutation; Deafness.
FT TRANSMEM 45 65 POTENTIAL.
FT TRANSMEM 98 118 POTENTIAL.
FT TRANSMEM 132 152 POTENTIAL.
FT TRANSMEM 173 193 POTENTIAL.
FT TRANSMEM 238 258 POTENTIAL.
FT TRANSMEM 298 318 POTENTIAL.
FT TRANSMEM 276 276 W -> S (IN DNAA2).
FT VARIANT /FTId=VAR_008726.
FT VARIANT G -> C (IN DNAA2; LOSS OF POTASSIUM
SELECTIVITY OF THE PORE).
FT VARIANT /FTId=VAR_008727.
FT VARIANT G -> S (IN DNAA2).
FT VARIANT /FTId=VAR_001547.
FT VARIANT G -> S (IN DNAA2).
FT VARIANT /FTId=VAR_008728.
SQ SEQUENCE 695 AA; 77091 MW; A58737BD845E1A3A CRC64;

Query Match 5.3%; Score 20; DB 1; Length 695;
Best Local Similarity 100.0%; Pred. No. 2.4e-12;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 303 ISFPALPAGILGSGFALKVQ 322
DB 308 ISFPALPAGILGSGFALKVQ 327
|||||
RESULT 4
C1Q3_HUMAN STANDARD; PRT; 872 AA.
AC Q43525;
DT 15-JUL-1999 (Rel. 38, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DT 01-OCT-2000 (Rel. 40, Last annotation update)
DE VOLTAGE-GATED POTASSIUM CHANNEL PROTEIN KQT-LIKE 3.
GN KCNQ3
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=9087323; PubMed=9872318;
RA Schroeder B.C., Kubisch C., Stein V., Jentsch T.J.;
RT "Moderate loss of function of cyclic-AMP-modulated KCNQ2/KCNQ3 K+
channels causes epilepsy.";
RL Nature 396:687-690 (1998).
RN [2]
RP SEQUENCE OF 48-872 FROM N.A., AND VARIANT EBN2 VAL-310.
RX MEDLINE=98085869; PubMed=9425900;
RA Charlier C., Singh N.A., Ryan S.G., Lewis T.B., Reus B.E., Leach R.J.,
RA Leppert M.;
RT "A pore mutation in a novel KQT-like potassium channel gene in an
idiopathic epilepsy family.";
RL Nat. Genet. 18:53-55 (1998).
CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.
CC -1- DOMAIN: THE SEGMENT S4 IS PROBABLY THE VOLTAGE-SENSOR AND IS
CC CHARACTERIZED BY A SERIES OF POSITIVELY CHARGED AMINO ACIDS AT
CC EVERY THIRD POSITION (BY SIMILARITY).
CC -1- DISEASE: DEFECTS IN KCNQ3 ARE THE CAUSE OF BENIGN FAMILIAL
CC NEONATAL CONVULSIONS TYPE 2 (BFNC2); ALSO KNOWN AS EPILEPSY,

CC BENIGN NEONATAL TYPE 2 (EBN2) BFNC2 IS AN AUTOSOMAL-DOMINANT
CC DISEASE.
CC -1- SIMILARITY: THIS CHANNEL PROTEIN BELONGS TO THE DELAYED RECTIFIER
CC CLASS. KQT SUBFAMILY.
CC
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CC
CC EMBL; AF071491; AAC96101.1; -.
DR EMBL; AF071478; AAC96101.1; JOINED.
DR EMBL; AF071479; AAC96101.1; JOINED.
DR EMBL; AF071480; AAC96101.1; JOINED.
DR EMBL; AF071481; AAC96101.1; JOINED.
DR EMBL; AF071482; AAC96101.1; JOINED.
DR EMBL; AF071483; AAC96101.1; JOINED.
DR EMBL; AF071484; AAC96101.1; JOINED.
DR EMBL; AF071485; AAC96101.1; JOINED.
DR EMBL; AF071486; AAC96101.1; JOINED.
DR EMBL; AF071487; AAC96101.1; JOINED.
DR EMBL; AF071488; AAC96101.1; JOINED.
DR EMBL; AF071489; AAC96101.1; JOINED.
DR EMBL; AF071490; AAC96101.1; JOINED.
DR EMBL; AF033347; AAB97314.1; -.
DR MIM; 602232; -.
DR MIM; 121201; -.
DR InterPro; IPR000636; -.
DR InterPro; IPR003091; -.
DR Pfam; PF00520; Ion_trans; 1.
DR PRINTS; PR00169; KCHANNEL.
KW Ionic channel; Transmembrane; Ion transport; Voltage-gated channel;
KW Multigene family; Disease mutation.
FT TRANSMEM 122 142 POTENTIAL.
FT TRANSMEM 153 173 POTENTIAL.
FT TRANSMEM 197 217 POTENTIAL.
FT TRANSMEM 262 282 POTENTIAL.
FT TRANSMEM 300 320 POTENTIAL.
FT TRANSMEM 331 351 POTENTIAL.
FT VARIANT 310 310 G -> V (IN BFNC2).
FT /FTId=VAR_001546.
SQ SEQUENCE 872 AA; 96742 MW; BB79C69EE8591A84 CRC64;

Query Match 3.5%; Score 13; DB 1; Length 872;
Best Local Similarity 100.0%; Pred. No. 4.3e-05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 304 SFPAIPAGILGSG 316
DB 342 SFPAIPAGILGSG 354
|||||
RESULT 5
YJCC_ECOLI STANDARD; PRT; 528 AA.
ID YJCC_ECOLI
AC P32701; P76789;
DT 01-OCT-1993 (Rel. 27, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE HYPOTHETICAL 60.8 KDA PROTEIN IN SSB-SOX5 INTERGENIC REGION (0528).
OS Escherichia coli.
OC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
OC Escherichia.
OX NCBI_TaxID=562;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=K12 / MG1655;
RX MEDLINE=94089392; PubMed=8265357;

RT "Molecular genetics of the long QT syndrome: two novel mutations of
RT the KVLQT1 gene and phenotypic expression of the mutant gene in a
RT large kindred.";
RL Hum. Mutat. 11:158-165(1998).
RN [16]
RN VARIANT LQT1 PHE-339 DEL.
RX MEDLINE=9836456; PubMed=9702906;
RA Ackerman M.J., Schroeder J.J., Berry R., Schaid D.J., Porter C.-B.J.,
RA Michels V.V., Thibodeau S.N.;
RT "A novel mutation in KVLQT1 is the molecular basis of inherited long
RT QT syndrome in a near-drowning patient's family.";
RL Pediatr. Res. 44:148-153(1998).
RN [17]
RN VARIANT LQT1 THR-525.
RX MEDLINE=99415293; PubMed=10482963;
RA Larsen L.A., Fosdal I., Andersen P.S., Kanter J.K., Vuust J.,
RA Wetrell G., Christiansen M.;
RT "Recessive Romano-Ward syndrome associated with compound
RT heterozygosity for two mutations in the KVLQT1 gene.";
RL Eur. J. Hum. Genet. 7:724-728(1999).
RN [18]
RN VARIANTS LQT1 S-184; R-189; S-314; S-315; R-345; P-373 AND R-392.
RX MEDLINE=99235550; PubMed=10220144;
RA Jongbloed R.J.E., Wilde A.A.M., Geelen J.L.M.C., Doevendans P.,
RA Schaap C., van Langen I., van Tintelen J.P., Cobben J.M.,
RA Beaufort-Krol G.C.M., Geraedts J.P.M., Smeets H.J.M.;
RT "Novel KCNQ1 and HERG missense mutations in Dutch long-QT families.";
RL Hum. Mutat. 13:301-310(1999).
RN [19]
RN VARIANT LQT1 CYS-157.
RX MEDLINE=99235552; PubMed=10220146;
RA Larsen L.A., Christiansen M., Vuust J., Andersen P.S.;
RT "High-throughput single-strand conformation polymorphism analysis by
RT automated capillary electrophoresis: robust multiplex analysis and
RT pattern-based identification of allelic variants.";
RL Hum. Mutat. 13:318-327(1999).
RN [20]
RN VARIANTS LQT1 GLN-190; HIS-243; TRP-533 AND TRP-539.
RX MEDLINE=20192867; PubMed=10728423;
RA Chouabe C., Neyroud N., Richard P., Denjoy I., Hainque B., Roney G.,
RA Drici M.D., Guicheney P., Barhanin J.;
RT "Novel mutations in KVLQT1 that affect Iks activation through
RT interactions with Isk.";
RL Cardiovasc. Res. 45:971-980(2000).
RN [21]
RN VARIANTS LQT1.
RX MEDLINE=20432616; PubMed=10973849;
RA Splawski I., Shen J., Timothy K.W., Lehmann M.H., Priori S.,
RA Robinson J.L., Moss A.J., Schwartz P.J., Towbin J.A., Vincent G.M.,
RA Keating M.F.;
RT "Spectrum of mutations in long-QT syndrome genes. KVLQT1, HERG, SCN5A,
RT KCNE1, and KCNE2.";
RL Circulation 102:1178-1185(2000).
RN [22]
RN FUNCTION: PROBABLY IMPORTANT IN CARDIAC REPOLARIZATION. ASSOCIATES
CC WITH KCNE1 (MINK) TO FORM THE I(KS) CARDIAC POTASSIUM CURRENT.
CC ELICITS A RAPIDLY ACTIVATING, K(+) SELECTIVE OUTWARD CURRENT.
CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.
CC -1- ALTERNATIVE PRODUCTS: A NUMBER OF FORMS ARE PRODUCED BY
CC ALTERNATIVE SPLICING. KVLQT1 IS A TRUNCATED ISOFORM THAT IS
CC NONFUNCTIONAL ALONE BUT MODULATORY WHEN COEXPRESSED WITH THE FULL-
CC LENGTH ISOFORM.
CC -1- DOMAIN: THE SEGMENT S4 IS PROBABLY THE VOLTAGE-SENSOR AND IS
CC CHARACTERIZED BY A SERIES OF POSITIVELY CHARGED AMINO ACIDS AT
CC EVERY THIRD POSITION.
CC -1- DISEASE: DEFECTS IN KCNQ1 IS THE CAUSE OF LONG QT SYNDROME TYPE 1
CC (LQT1 OR LOTS). LQT1 IS A CONGENITAL HEART DISEASE WITH FREQUENT
CC FAMILIAL TRANSMISSION AND IS CHARACTERIZED BY A PROLONGED QT
CC INTERVAL IN THE ELECTROCARDIOGRAM WHICH CAUSES ABNORMAL

Query Match

18.4%; Score 69; DB 1; Length 676;

Best Local Similarity 100.0%; Pred. No. 1.5e-62;

Matches 69; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY .185 GQVFATSAIRGIRFLQILRLMLHVDROGGTWRLLGSGVVFHROELITTYIGFLGIFSSY 244
DB .219 GQVFATSAIRGIRFLQILRLMLHVDROGGTWRLLGSGVVFHROELITTYIGFLGIFSSY 278
QY 245 FVYLAEKDA 253
DB 279 FVYLAEKDA 287
RESULT 3
C1Q4_HUMAN
ID C1Q4_HUMAN STANDARD; PRT; 695 AA.
AC P56696; O96025;
DT 15-JUL-1999 (Rel. 38, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DE 01-OCT-2000 (Rel. 40, Last annotation update)
DE VOLTAGE-GATED POTASSIUM CHANNEL PROTEIN KQT-LIKE 4.
GN KCNQ4.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RN SEQUENCE FROM N.A., AND VARIANT DFNA2 SER-285.
RX MEDLINE=99148276; PubMed=10025409;
RA Kubisch C., Schroeder B.C., Friedrich T., Luetjohann B.,
RA El-Amraoui A., Marlin S., Petit C., Jentsch T.J.;
RT "KCNQ4, a novel potassium channel expressed in sensory outer hair
RT cells, is mutated in dominant deafness.";
RL Cell 96:437-446(1999).
RN [2]
RN VARIANTS DFNA2 SER-276; CYS-285 AND SER-321.
RX MEDLINE=99299248; PubMed=10369879;
RA Coucke P.J., Van Hauwe P., Kelley P.M., Kunst H., Schattelman I.,
RA Van Velzen D., Meyers J., Ensink R.J., Verstreken M., Declau F.,
RA Marres H., Kastury K., Bhasin S., McGuirt W.T., Smith R.J.H.,
RA Cremers C.W.R.J., Van de Heyning P., Willems P.J., Smith S.D.,
RA Van Camp G.;
RT "Mutations in the KCNQ4 gene are responsible for autosomal dominant
RT deafness in four DFNA2 families.";
RL Hum. Mol. Genet. 8:1321-1328(1999).
CC -1- FUNCTION: MAY BE RESPONSIBLE FOR POTASSIUM IONS AFTER STIMULATION
CC OF THE HAIR CELL.
CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.
CC -1- TISSUE SPECIFICITY: EXPRESSED IN THE OUTER, BUT NOT THE INNER,
CC SENSORY HAIR CELLS OF THE COCHLEA.
CC -1- DOMAIN: THE SEGMENT S4 IS PROBABLY THE VOLTAGE-SENSOR AND IS
CC CHARACTERIZED BY A SERIES OF POSITIVELY CHARGED AMINO ACIDS AT
CC EVERY THIRD POSITION (BY SIMILARITY).
CC -1- DISEASE: DEFECTS IN KCNQ4 ARE A CAUSE OF AUTOSOMAL DOMINANT
CC NONSYNDROMIC SENSORINEURAL DEAFNESS TYPE 2 (DFNA2).
CC -1- SIMILARITY: THIS CHANNEL PROTEIN BELONGS TO THE DELAYED RECTIFIER
CC CLASS. KQT SUBFAMILY.
CC -1- This SWISS-PROT entry is copyright. It is produced through a collaboration
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CC -----
CC EMBL; AF105202; AAD14680.1;
CC EMBL; AF105216; AAD14681.1;
CC EMBL; AF105203; AAD14681.1; JOINED.
CC EMBL; AF105204; AAD14681.1; JOINED.
CC EMBL; AF105205; AAD14681.1; JOINED.
CC EMBL; AF105206; AAD14681.1; JOINED.
CC EMBL; AF105207; AAD14681.1; JOINED.
CC EMBL; AF105208; AAD14681.1; JOINED.
CC EMBL; AF105209; AAD14681.1; JOINED.
CC EMBL; AF105210; AAD14681.1; JOINED.
CC EMBL; AF105211; AAD14681.1; JOINED.

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OM protein - protein search, using sw model

Run on: November 2, 2001, 12:02:04 ; Search time 27.64 Seconds
(without alignments)
465.994 Million cell updates/sec

Title: US-09-135-010A-113
Perfect score: 376
Sequence: 1 MNENAINSLYEAIPLPDGGS.....TWKIYIKQSRNHHIMSPSP 376

Scoring table: OLIGO
Gapop 60.0 , Gapext 60.0

Searched: 93435 seqs, 34255486 residues

Word size : 4

Total number of hits satisfying chosen parameters: 60339

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : SwissProt_39:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	71	18.9	604	1	P97414 mus musculus
2	69	18.4	676	1	P51787 homo sapien
3	20	5.3	695	1	P56696 homo sapien
4	13	3.5	872	1	O43525 homo sapien
5	10	2.7	528	1	P32701 escherichia
6	7	1.9	119	1	O95298 homo sapien
7	7	1.9	173	1	Q13185 homo sapien
8	7	1.9	176	1	Q24808 entamoeba d
9	7	1.9	177	1	Q24810 entamoeba h
10	7	1.9	241	1	O60274 methanococc
11	7	1.9	244	1	P37813 bacillus su
12	7	1.9	256	1	P03565 cassava lat
13	7	1.9	261	1	P03698 bacterioph
14	7	1.9	268	1	O49402 mycoplasma
15	7	1.9	332	1	P44984 caenophilus
16	7	1.9	335	1	P34410 haemorrhabi
17	7	1.9	387	1	P45550 escherichia
18	7	1.9	395	1	P53554 bacillus su
19	7	1.9	460	1	Q36458 ornithorhyn
20	7	1.9	474	1	P15581 paramecium
21	7	1.9	485	1	Q00270 hepatitis e
22	7	1.9	486	1	P08465 saccharomyc
23	7	1.9	624	1	O04130 arabidopsis
24	7	1.9	660	1	P29326 hepatitis e
25	7	1.9	660	1	Q04611 hepatitis e
26	7	1.9	660	1	P33426 hepatitis e
27	7	1.9	669	1	P78588 candida alb
28	7	1.9	707	1	Q07266 rattus norv
29	7	1.9	722	1	Q42667 citrus limo
30	7	1.9	752	1	P28840 rattus norv
31	7	1.9	875	1	P41812 saccharomyc
32	7	1.9	926	1	O74756 schizosacch
33	7	1.9	1442	1	Q10570 homo sapien

Query Match 18.9% Score 71; DB 1; Length 604;

34 1.9 1444 1 CPSA_BOVIN--
35 7 1.9 1447 1 SGSL_YEAST
36 6 1.6 52 1 CRAB_TRASC
37 6 1.6 57 1 V3A_IBVM
38 6 1.6 57 1 V3A_IBVP3
39 6 1.6 57 1 V3A_IBVU5
40 6 1.6 58 1 V3A_IBVB
41 6 1.6 63 1 ITHV_HIRMA
42 6 1.6 66 1 CYT_SOLTU
43 6 1.6 66 1 GYPA_AMOPE
44 6 1.6 71 1 YVFE_VACCC
45 6 1.6 72 1 HTF_BLADI

ALIGNMENTS

RESULT 1
C1Q1_MOUSE STANDARD; PRT; 604 AA.
AC P97414;
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE VOLTAGE-GATED POTASSIUM CHANNEL PROTEIN KQT-LIKE 1 (KV1.9).
GN KCNQ1 OR KCNA9 OR KVLQT1.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=97055937; PubMed=8900282;
RA Barhanin J., Lesage F., Guillemare E., Fink M., Lazdunski M.,
RA Ramey G.;
RT "K(V)LQT1 and Isk (minK) proteins associate to form the I(Ks)
RT potassium current".
RL Nature 384:78-80(1996).
CC - FUNCTION: PROBABLY IMPORTANT IN CARDIAC REPOLARIZATION. ASSOCIATES
CC WITH KCNE1 (MINK) TO FORM THE I(KS) CARDIAC POTASSIUM CURRENT.
CC ELICITS A RAPIDLY ACTIVATING, K(+) SELECTIVE OUTWARD CURRENT.
CC - SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.
CC - DOMAIN: THE SEGMENT S4 IS PROBABLY THE VOLTAGE-SENSOR AND IS
CC CHARACTERIZED BY A SERIES OF POSITIVELY CHARGED AMINO ACIDS AT
CC EVERY THIRD POSITION.
CC - SIMILARITY: THIS CHANNEL PROTEIN BELONGS TO THE DELAYED RECTIFIER
CC CLASS. KQT SUBFAMILY.
CC
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CC
CC EMBL: U70068; AAB36518.1; -
CC MCD; MGI:108083; Kcnql.
CC InterPro: IPR000636; -
CC Pfam: PF00520; Ion.Trans. 1.
CC Ionic channel; Transmembrane; Ion transport; Voltage-gated channel;
CC Glycoprotein; Multigene family; Phosphorylation.
FT TRANSMEM 57 77 SEGMENT S1 (POTENTIAL).
FT TRANSMEM 83 103 SEGMENT S2 (POTENTIAL).
FT TRANSMEM 132 152 SEGMENT S3 (POTENTIAL).
FT TRANSMEM 197 217 SEGMENT S4 (POTENTIAL).
FT TRANSMEM 231 251 SEGMENT S5 (POTENTIAL).
FT TRANSMEM 263 283 SEGMENT S6 (POTENTIAL).
FT CARBOHYD 224 224 N-LINKED (GLCNAC...)(POTENTIAL).
SQ SEQUENCE 604 AA; 68070 MW; 0E3579910F1CB697 CRC64;